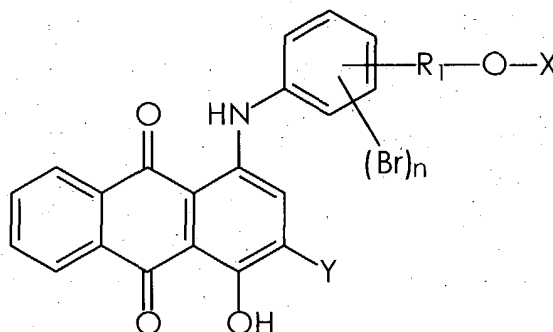


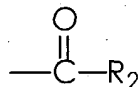
PHASE CHANGE INKS CONTAINING COLORANT COMPOUNDS

Cross-reference is made to the following copending applications:

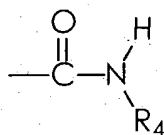
- 5 Copending Application U.S. Serial No. 10/260,146, filed September 27, 2002, entitled "Colorant Compounds," with the named inventors Jeffery H. Banning and C. Wayne Jaeger, the disclosure of which is totally incorporated herein by reference, discloses compounds of the formula



- 10 wherein Y is a hydrogen atom or a bromine atom, n is an integer of 0, 1, 2, 3, or 4, R_1 is an alkylene group or an arylalkylene group, and X is (a) a hydrogen atom, (b) a group of the formula

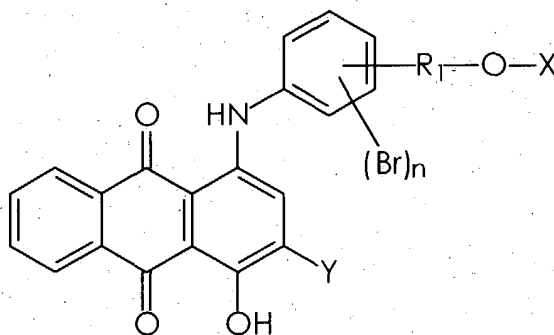


- 15 wherein R_2 is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group, (c) an alkyleneoxy, aryleneoxy, arylalkyleneoxy, or alkylaryleneoxy group, or (d) a group of the formula



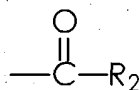
wherein R_4 is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group.

Copending Application U.S. Serial No. 10/260,376, filed
 5 September 27, 2002, entitled "Phase Change Inks," with the named
 inventors C. Wayne Jaeger and Jeffery H. Banning, the disclosure of
 which is totally incorporated herein by reference, discloses a phase
 change ink composition comprising a phase change ink carrier and a
 colorant compound of the formula

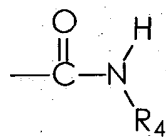


10

wherein Y is a hydrogen atom or a bromine atom, n is an integer of 0, 1,
 2, 3, or 4, R_1 is an alkylene group or an arylalkylene group, and X is (a) a
 hydrogen atom, (b) a group of the formula

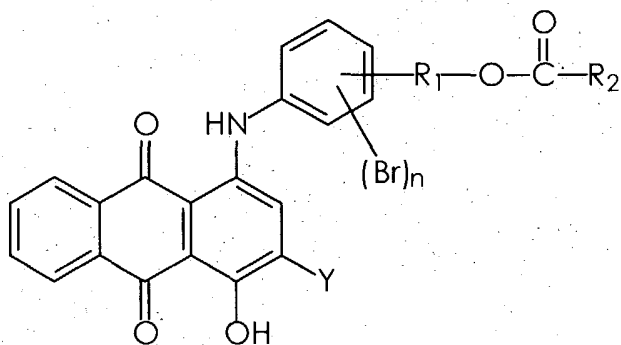


15 wherein R_2 is an alkyl group, an aryl group, an arylalkyl group, or an
 alkylaryl group, (c) an alkyleneoxy, aryleneoxy, arylalkyleneoxy, or
 alkylaryleneoxy group, or (d) a group of the formula

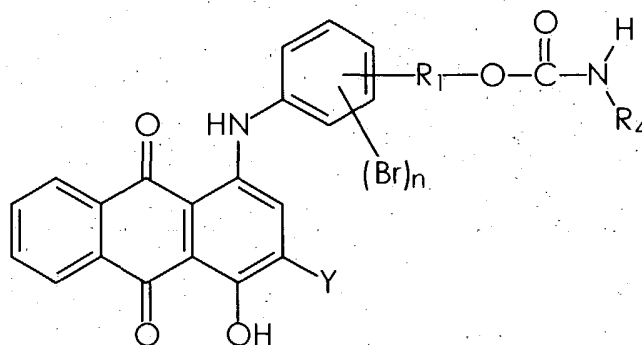


wherein R₄ is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group.

Copending Application U.S. Serial No. 10/260,379, filed September 27, 2002, entitled "Methods for Making Colorant Compounds," with the named inventors C. Wayne Jaeger and Jeffery H. Banning, the disclosure of which is totally incorporated herein by reference, discloses a process for preparing a colorant of the formula

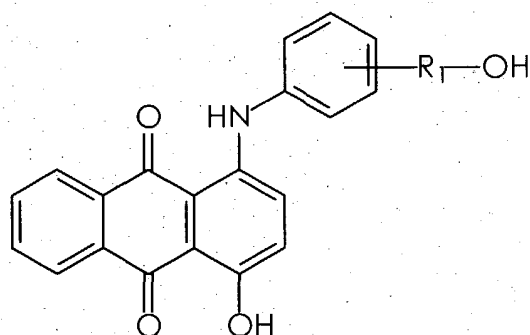


10 or

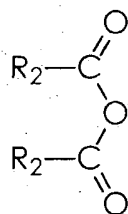


wherein Y is a hydrogen atom or a bromine atom, n is an integer of 0, 1, 2, 3, or 4, R₁ is an alkylene group or an arylalkylene group, R₂ is an alkyl

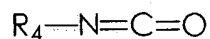
group, an aryl group, an arylalkyl group, or an alkylaryl group, and R_4 is an alkyl group, an aryl group, an arylalkyl group, or an alkylaryl group, can be prepared by a process which comprises (a) preparing a first reaction mixture by admixing (1) leucoquinizarin and, optionally,
 5 quinizarin, (2) an aminobenzene substituted with an alcohol group of the formula $-R_1-OH$, (3) boric acid, and (4) an optional solvent, and heating the first reaction mixture to prepare an alcohol-substituted colorant of the formula



10 followed by (b) converting the colorant thus prepared to either (i) an ester-substituted colorant by reaction with an esterification compound which is either (A) an anhydride of the formula

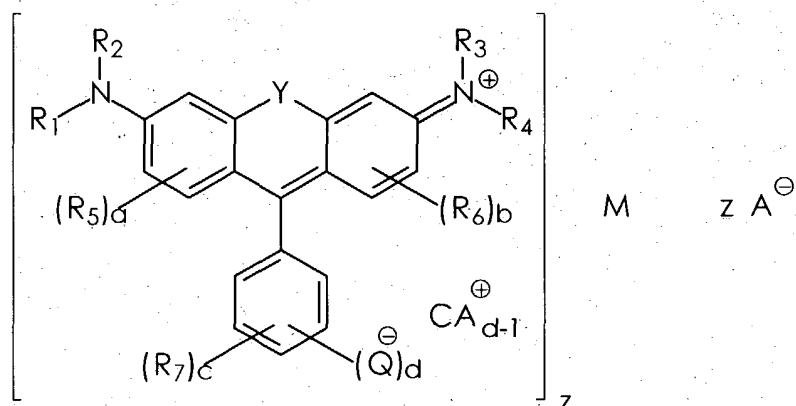


or (B) an acid of the formula R_2COOH in the presence of an optional
 15 esterification catalyst, or (ii) a urethane-substituted colorant by reaction with an isocyanate compound of the formula

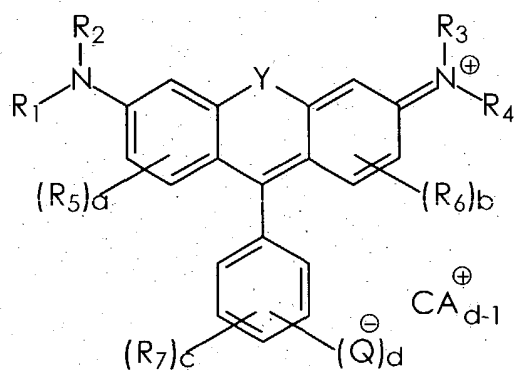


and (c) brominating the colorant thus prepared, wherein either conversion to ester or urethane can be performed before bromination or bromination can be performed before conversion to ester or urethane.

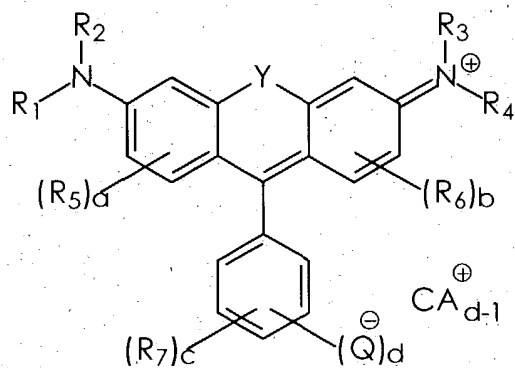
- 5 Copending Application U.S. Serial No. (not yet assigned; Attorney Docket Number D/A3152), filed concurrently herewith, entitled "Colorant Compounds," with the named inventors Jeffery H. Banning, Bo Wu, James M. Duff, Wolfgang G. Wedler, Jule W. Thomas, and Randall R. Bridgeman, the disclosure of which is totally incorporated herein by
10 reference, discloses compounds of the formula



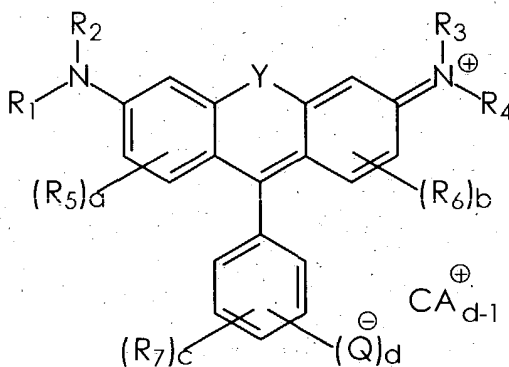
wherein M is either (1) a metal ion having a positive charge of +y wherein y is an integer which is at least 2, said metal ion being capable of forming a compound with at least two



chromogen moieties, or (2) a metal-containing moiety capable of forming a compound with at least two

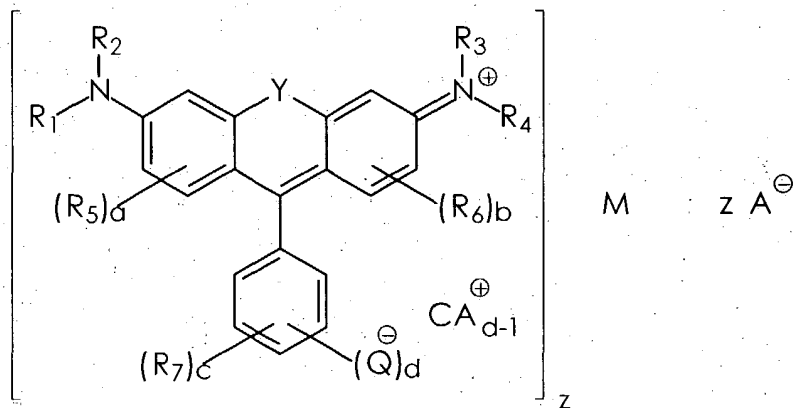


5 chromogen moieties, z is an integer representing the number of

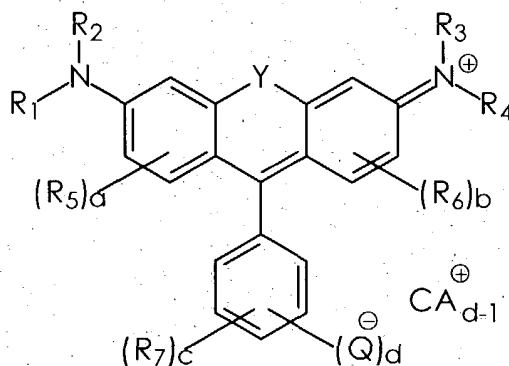


chromogen moieties associated with the metal and is at least 2, and R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , a , b , c , d , Y , Q , A , and CA are as defined therein.

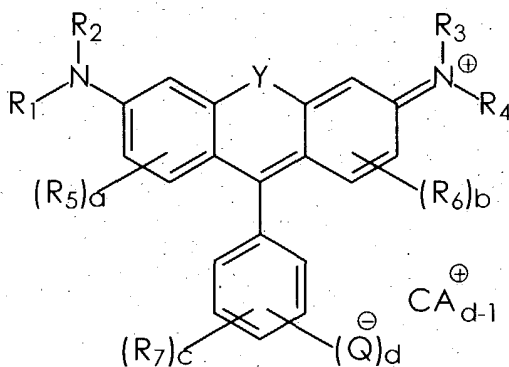
- Copending Application U.S. Serial No. (not yet assigned;
 5 Attorney Docket Number D/A3152Q), filed concurrently herewith,
 entitled "Phase Change Inks Containing Colorant Compounds," with the
 named inventors Bo Wu, Jeffery H. Banning, James M. Duff, Wolfgang
 G. Wedler, Jule W. Thomas, and Randall R. Bridgeman, the disclosure of
 which is totally incorporated herein by reference, discloses a phase
 10 change ink composition comprising a phase change ink carrier and a
 colorant compound of the formula



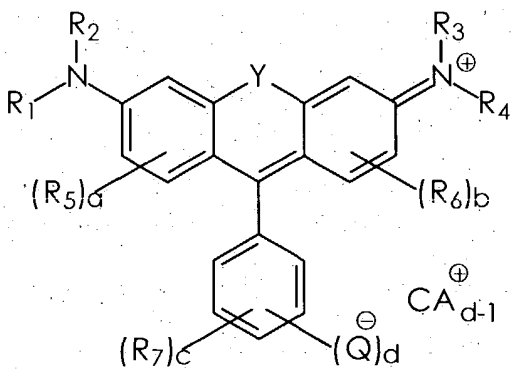
wherein M is either (1) a metal ion having a positive charge of +y wherein y is an integer which is at least 2, said metal ion being capable of forming a compound with at least two



5 chromogen moieties, or (2) a metal-containing moiety capable of forming a compound with at least two

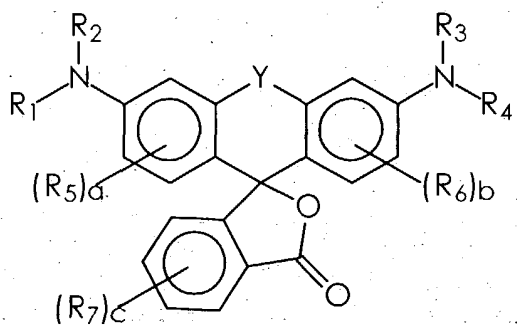


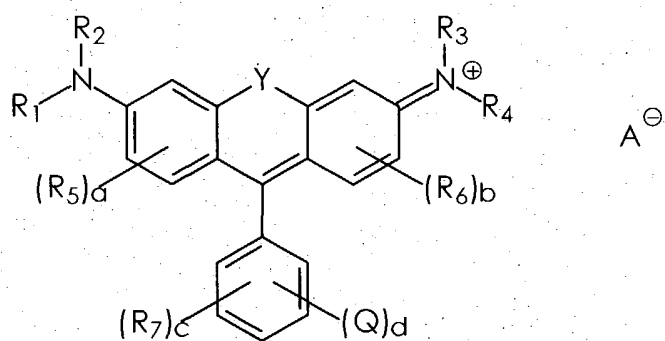
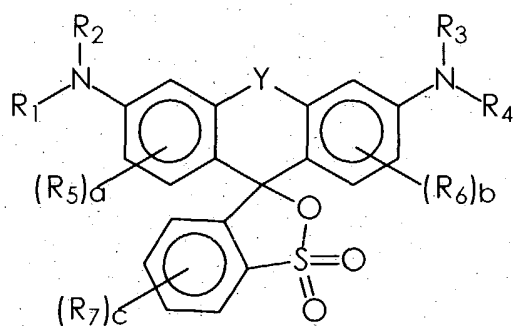
chromogen moieties, z is an integer representing the number of



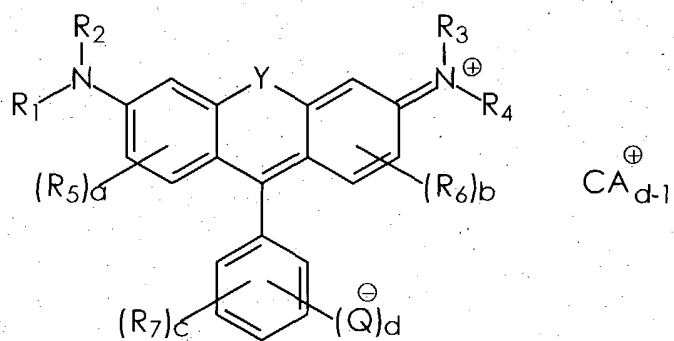
chromogen moieties associated with the metal and is at least 2, and R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , a , b , c , d , Y , Q , A , and CA are as defined therein.

- Copending Application U.S. Serial No. (not yet assigned;
 5 Attorney Docket Number D/A3153), filed concurrently herewith, entitled
 "Colorant Compounds," with the named inventors Jeffery H. Banning, Bo
 Wu, James M. Duff, Wolfgang G. Wedler, and Donald R. Titterington, the
 disclosure of which is totally incorporated herein by reference, discloses
 compounds of the formulae





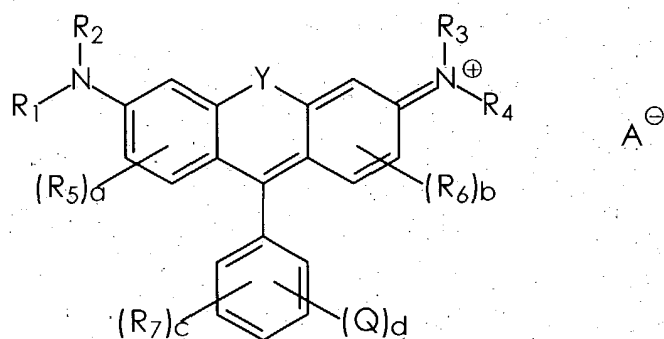
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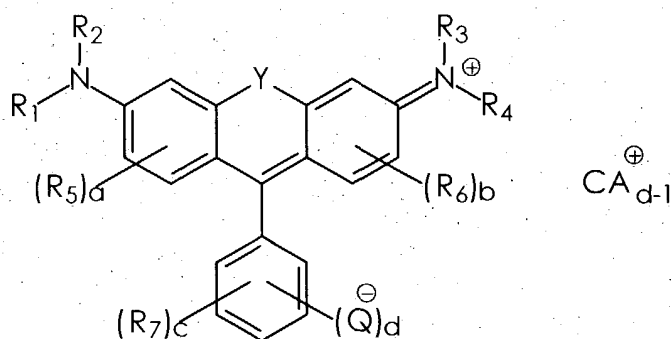
- 5 wherein R₁, R₂, R₃, R₄, R₅, R₆, R₇, a, b, c, d, Y, Q, Q⁻, A, and CA are as defined therein.

BACKGROUND

The present invention is directed to phase change inks. More specifically, the present invention is directed to hot melt or phase change inks containing specific colorant compounds. One embodiment of the present invention is directed to a phase change ink carrier and a colorant compound of the formula

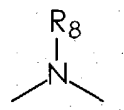
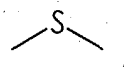
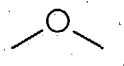
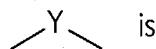


or

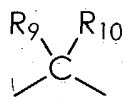


- 10 wherein R_1 , R_2 , R_3 , and R_4 each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group, (iii) an aryl group, (iv) an arylalkyl group, or (v) an alkylaryl group, wherein R_1 and R_2 can be joined together to form a ring, wherein R_3 and R_4 can be joined together to form a ring, and wherein R_1 , R_2 , R_3 , and R_4 can each be joined to a

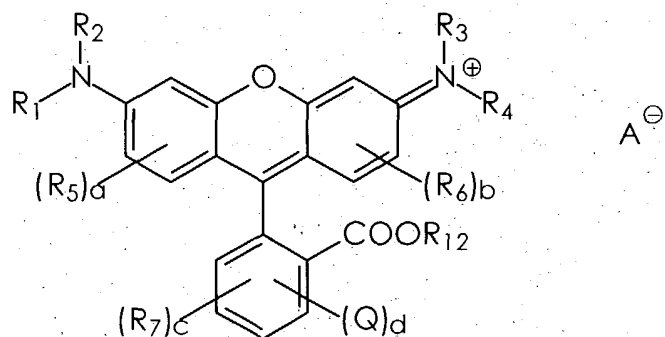
phenyl ring in the central structure, a and b each, independently of the others, is an integer which is 0, 1, 2, or 3, c is an integer which is 0, 1, 2, 3, or 4, each R₅, R₆, and R₇, independently of the others, is (i) an alkyl group, (ii) an aryl group, (iii) an arylalkyl group, (iv) an alkylaryl group, (v) a halogen atom, (vi) an ester group, (vii) an amide group, (viii) a sulfone group, (ix) an amine group or ammonium group, (x) a nitrile group, (xi) a nitro group, (xii) a hydroxy group, (xiii) a cyano group, (xiv) a pyridine or pyridinium group, (xv) an ether group, (xvi) an aldehyde group, (xvii) a ketone group, (xviii) a carbonyl group, (xix) a thiocarbonyl group, (xx) a sulfate group, (xxi) a sulfide group, (xxii) a sulfoxide group, (xxiii) a phosphine or phosphonium group, (xxiv) a phosphate group, (xxv) a mercapto group, (xxvi) a nitroso group, (xxvii) an acyl group, (xxviii) an acid anhydride group, (xxix) an azide group, (xxx) an azo group, (xxxi) a cyanato group, (xxxii) an isocyanato group, (xxxiii) a thiocyanato group, (xxxiv) an isothiocyanato group, (xxxv) a urethane group, or (xxxvi) a urea group, wherein R₅, R₆, and R₇ can each be joined to a phenyl ring in the central structure,



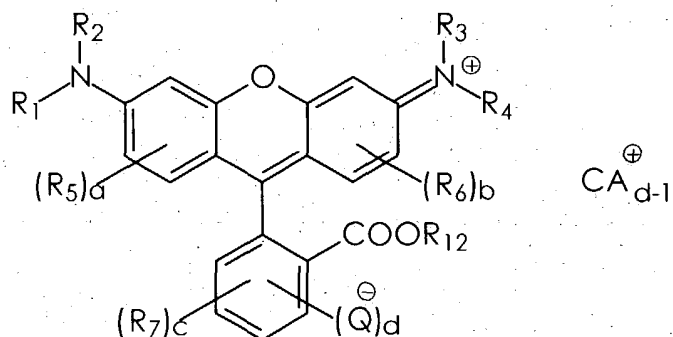
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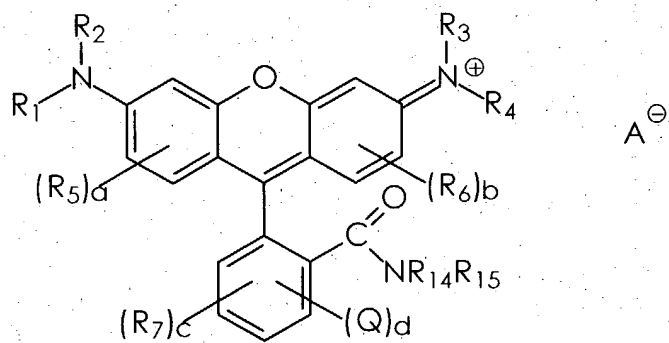
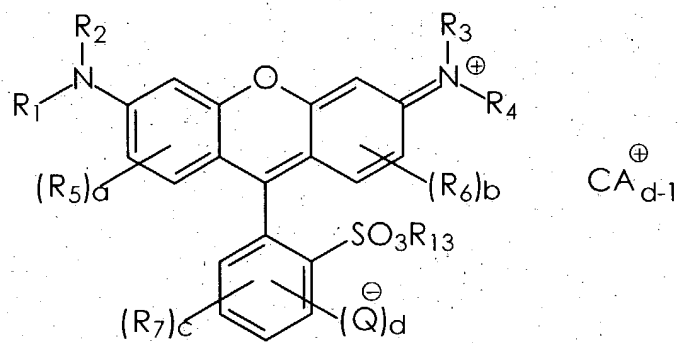
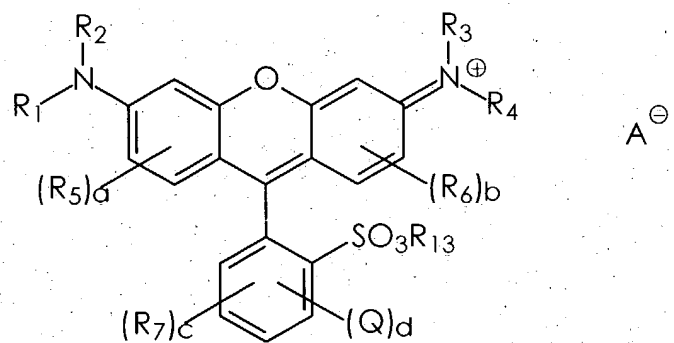


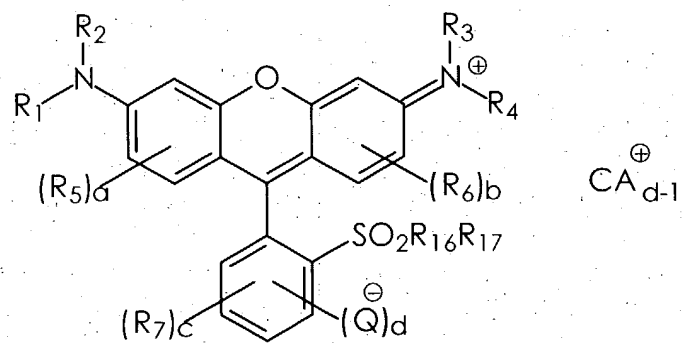
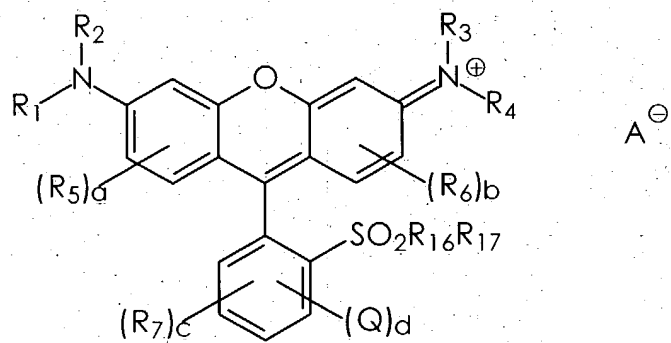
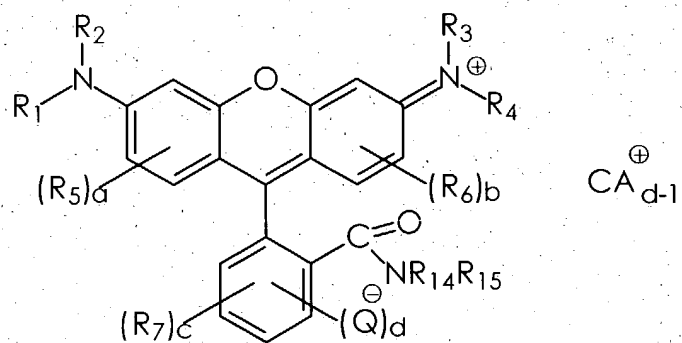
R_8 , R_9 , and R_{10} each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group, (iii) an aryl group, (iv) an arylalkyl group, or (v) an alkylaryl group, provided that the number of carbon atoms in $R_1+R_2+R_3+R_4+R_5+R_6+R_7+R_8+R_9+R_{10}$ is at least about 16, Q is a $COOH$ group or a SO_3H group, Q^- is a COO^- group or a SO_3^- group, d is an integer which is 1, 2, 3, 4, or 5, A is an anion, and CA is either a hydrogen atom or a cation associated with all but one of the Q^- groups, provided that when the colorant compound is of the formula



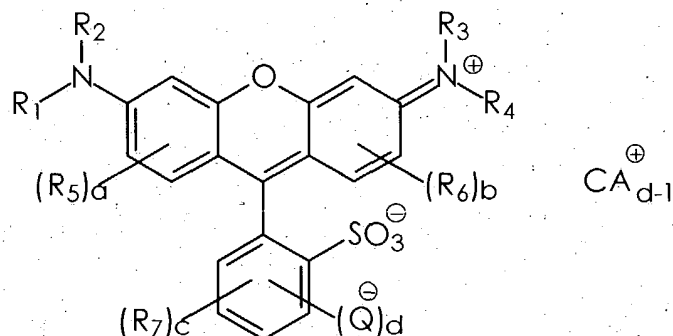
10







or



wherein R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , and R_7 each, independently of the other, is (i) an alkyl group, (ii) an aryl group, (iii) an arylalkyl group, or (iv) an alkylaryl group, (I) either (a) c is an integer which is 0, 1, 2, or 3, or (b) d is an integer which is 1, 2, 3, or 4, and (II) either (a) three of R_1 , R_2 , R_3 , and R_4 are hydrogen atoms; (b) only one of R_1 , R_2 , R_3 , and R_4 is a hydrogen atom; (c) R_1 and R_2 are both hydrogen atoms; (d) R_3 and R_4 are both hydrogen atoms; or (e) R_1 and R_3 are both hydrogen atoms and R_2 and R_4 are each, independently of the other, either alkyl groups or arylalkyl groups.

In general, phase change inks (sometimes referred to as "hot melt inks") are in the solid phase at ambient temperature, but exist in the liquid phase at the elevated operating temperature of an ink jet printing device. At the jet operating temperature, droplets of liquid ink are ejected from the printing device and, when the ink droplets contact the surface of the recording substrate, either directly or via an intermediate heated transfer belt or drum, they quickly solidify to form a predetermined pattern of solidified ink drops. Phase change inks have also been used in other printing technologies, such as gravure printing, as disclosed in, for example, U.S. Patent 5,496,879 and German Patent

Publications DE 4205636AL and DE 4205713AL, the disclosures of each of which are totally incorporated herein by reference.

Phase change inks for color printing typically comprise a phase change ink carrier composition which is combined with a phase change ink compatible colorant. In a specific embodiment, a series of colored phase change inks can be formed by combining ink carrier compositions with compatible subtractive primary colorants. The subtractive primary colored phase change inks can comprise four component dyes, namely, cyan, magenta, yellow and black, although the inks are not limited to these four colors. These subtractive primary colored inks can be formed by using a single dye or a mixture of dyes. For example, magenta can be obtained by using a mixture of Solvent Red Dyes or a composite black can be obtained by mixing several dyes. U.S. Patent 4,889,560, U.S. Patent 4,889,761, and U.S. Patent 5,372,852, the disclosures of each of which are totally incorporated herein by reference, teach that the subtractive primary colorants employed can comprise dyes from the classes of Color Index (C.I.) Solvent Dyes, Disperse Dyes, modified Acid and Direct Dyes, and Basic Dyes. The colorants can also include pigments, as disclosed in, for example, U.S. Patent 5,221,335, the disclosure of which is totally incorporated herein by reference. U.S. Patent 5,621,022, the disclosure of which is totally incorporated herein by reference, discloses the use of a specific class of polymeric dyes in phase change ink compositions.

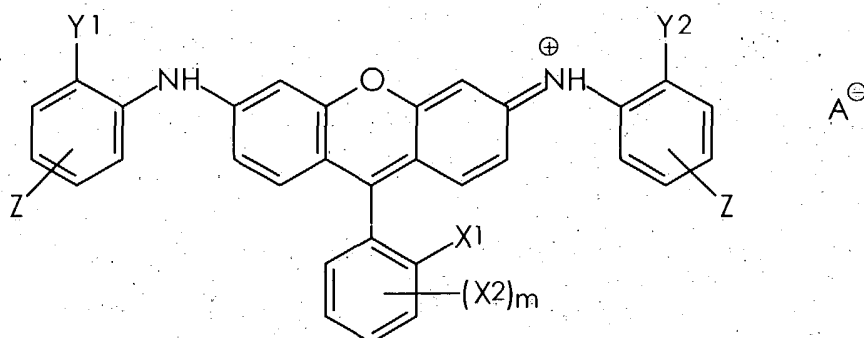
Phase change inks have also been used for applications such as postal marking, industrial marking, and labelling.

Phase change inks are desirable for ink jet printers because they remain in a solid phase at room temperature during shipping, long

term storage, and the like. In addition, the problems associated with nozzle clogging as a result of ink evaporation with liquid ink jet inks are largely eliminated, thereby improving the reliability of the ink jet printing. Further, in phase change ink jet printers wherein the ink droplets are applied directly onto the final recording substrate (for example, paper, transparency material, and the like), the droplets solidify immediately upon contact with the substrate, so that migration of ink along the printing medium is prevented and dot quality is improved.

Compositions suitable for use as phase change ink carrier compositions are known. Some representative examples of references disclosing such materials include U.S. Patent 3,653,932, U.S. Patent 4,390,369, U.S. Patent 4,484,948, U.S. Patent 4,684,956, U.S. Patent 4,851,045, U.S. Patent 4,889,560, U.S. Patent 5,006,170, U.S. Patent 5,151,120, U.S. Patent 5,372,852, U.S. Patent 5,496,879, European Patent Publication 0187352, European Patent Publication 0206286, German Patent Publication DE 4205636AL, German Patent Publication DE 4205713AL, and PCT Patent Application WO 94/04619, the disclosures of each of which are totally incorporated herein by reference. Suitable carrier materials can include paraffins, microcrystalline waxes, polyethylene waxes, ester waxes, fatty acids and other waxy materials, fatty amide containing materials, sulfonamide materials, resinous materials made from different natural sources (tall oil rosins and rosin esters, for example), and many synthetic resins, oligomers, polymers, and copolymers.

British Patent Publication GB 2 311 075 (Gregory et al.), the disclosure of which is totally incorporated herein by reference, discloses a compound of the formula



wherein X¹ is an ester group or an amide group (such as of a carboxylic or sulfonic acid) or a fatty amine salt of a sulfonic acid, each X² independently is a substituent, m has a value of from 0 to 2, Y¹ and Y² are each independently H, alkyl, or halo, each Z independently is an ester or amide group, and A⁻ is an anion. The compound is useful as a colorant for toners, D2T2 printing, plastics, polyesters, nylons, and inks, especially ink jet or hot melt inks.

"Rhodamine Dyes and Related Compounds. XV.

- 10 Rhodamine Dyes with Hydroaromatic and Polymethylene Radicals," I. S. Ioffe et al., *Zh. Organ. Khim.* (1965), 1(3), 584-6, the disclosure of which is totally incorporated herein by reference, discloses a process wherein heating dichlorofluoran with ZnCl₂-ZnO and the appropriate amine for 3 hours at 220° followed by treatment with aqueous HCl gave
- 15 N,N'-dicyclohexylrhodamine-HCl, m. 180-5°,
 N,N'-di(tetramethylene)rhodamine-HCl, decompd. 240°,
 N,N'-di(pentamethylene)rhodamine-HCl, m. 205-10°,
 N,N'-di(hexamethylene)rhodamine-HCl, decompd. 175°. These dyes gave yellow or orange fluorescence and their spectra were given.

- 20 "Rhodamine Dyes and Related Compounds. XI. Aryl- and Alkylrhodamines Containing Carboxyl Groups," I. S. Ioffe et al., *Zh.*

Obsch. Khim. (1964), 34(6), 2041-4, the disclosure of which is totally incorporated herein by reference, discloses a process wherein heating aminobenzoic acids with 3,6-dichlorofluoran in the presence of ZnCl_2 for 6 hours at $24-50^\circ$ gave after an aqueous treatment:

5 N,N'-bis(o-carboxyphenyl)rhodamine-HCl; m-isomer-HCl; and p-isomer-HCl. A similar reaction with HCl salts of glycine, α -alanine, or β -alanine gave: N,N'-bis(carboxymethyl)rhodamine-HCl; N,N'-bis(α -carboxyethyl)rhodamine-HCl; and N,N'-bis(β -carboxyethyl)rhodamine-HCl. The latter group showed

10 yellow-green fluorescence, lacking in the aryl derivatives. Spectra of the products are shown.

"Rhodamine Dyes and Related Compounds. X. Fluorescence of Solutions of Alkyl- and Arylalkylrhodamines," I. S. Ioffe et al., Zh. Obsch. Khim. (1964), 34(6), 2039-41, the disclosure of which is

15 totally incorporated herein by reference, discloses fluorescence spectra for the following rhodamines: N,N'-diethyl; N,N'-dibenzyl; N,N'-bis(β -phenylethyl); N,N'-bis(β -phenylisopropyl). In symmetrical substituted rhodamines, the entry of an alkyl or arylalkyl group into both amino residues resulted in the displacement of fluorescence max.

20 toward longer wavelengths, a similar displacement of absorption and an increase in the quantum yield of fluorescence. In unsymmetrical derivatives, an aryl group entering one of the amino groups shifted the spectra to a greater degree in the same direction and sharply reduced the quantum yield of fluorescence.

25 "Rhodamine Dyes and Related Compounds. IX. Rhodamine B Sulfonic Acids and their Derivatives," I. S. Ioffe et al., Zh. Obsch. Khim. (1964), 34(2), 640-44, the disclosure of which is totally

incorporated herein by reference, discloses that heating $m\text{-Et}_2\text{NC}_6\text{H}_4\text{OH}$ and $\text{K } \beta\text{-sulfophthalate}$ at 150° while concentrated H_2SO_4 was being added gave after 3 hours at $150\text{-}70^\circ$, followed by heating with H_2O 15 min., a residue of crude sulforhodamine, purified by solution in hot aqueous Na_2CO_3 and precipitation with AcOH . The mixed isomeric rhodamine sulfonic acids refluxed 3 hours with 30% AcOH , clarified, and cooled gave a first isomer with R_f 0.74 on paper in aqueous solution (pH 9) while the residue was the other isomer with R_f 0.98. The first isomer and PCl_5 gave the sulfonyl chloride, isolated as HCl salt, red solid (from $\text{CHCl}_3\text{-ligroine}$), which with NH_3 in CHCl_3 gave the sulfonamide, a violet powder. The two isomers and Rhodamine B had similar spectral characteristics. The two isomers probably contain the SO_3H group in the 4- and 5-positions of the Ph ring of Rhodamine B. Their absorption and fluorescence spectra are shown. Their solutions in CHCl_3 gave stronger fluorescence than those in Me_2CO .

"Rhodamine Dyes and Related Compounds. VIII. Amides of Sulforhodamine B Containing β -Hydroxyethyl and β -Chloroethyl Groups," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1963), 33(12), 3943-6, the disclosure of which is totally incorporated herein by reference, discloses that sulforhodamine B chloride heated 10-12 hours with $\text{HOCH}_2\text{CH}_2\text{NH}_2$ at $170\text{-}80^\circ$, then triturated with saturated NaCl gave, after solution in CHCl_3 and precipitation with petroleum ether, 80% red sulforhodamine B $\text{N}(\beta\text{-hydroxyethyl})\text{amide}$; similar reaction with $\text{HN}(\text{CH}_2\text{CH}_2\text{OH})_2$ gave 70% $\text{N,N-bis}(\beta\text{-hydroxyethyl})\text{amide}$, a bright red wax. These treated with SOCl_2 in CHCl_3 gave, respectively, $\text{N}(\beta\text{-chloroethyl})\text{amide}$, a brown powder, and $\text{N,N-bis}(\beta\text{-chloroethyl})\text{amide}$, a violet powder. Absorption

spectra of the amides are shown. The (hydroxyethyl)amides displayed strong orange fluorescence in solution.

"Rhodamine Dyes and Related Compounds. VII. (β -Phenylethyl)rhodamines," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1963), 33(4), 1089-92, the disclosure of which is totally incorporated herein by reference, discloses a process wherein heating dichlorofluoran with $\text{PhCH}_2\text{CH}_2\text{NH}_2$ or $\text{PhCH}_2\text{CH}(\text{Me})\text{NH}_2$ in the presence of ZnO and ZnCl_2 for 5-6 hours at 220° gave, after heating for 2 hours with aqueous HCl , 96-8% crude products which, after crystallization from alc. HCl , gave red, powdery $\text{N,N'$ -bis(β -phenylethyl)rhodamine- HCl , m. $172-5^\circ$, or $\text{N,N'$ -bis(α -methyl- β -phenylethyl)rhodamine- HCl , m. $175-8^\circ$; N -phenyl- N' -(β -phenylethyl)rhodamine- HCl , m. $162-6^\circ$, was prepared from $\text{PhCH}_2\text{CH}_2\text{NH}_2$ and 3'-chloro-6'-anilino-fluoran under the above conditions. Treated with alc. NaOH and quenched in H_2O , these hydrochlorides gave the free bases of the dyes as brown-red solids, which tended to form colloids in aqueous medium. The free bases m. $123-5^\circ$, decompd. 120° , and m. $164-8^\circ$, respectively. The ultraviolet and visible spectra of the dyes were similar to the spectra of dibenzylrhodamine, but had deeper color; strong fluorescence was shown by these dyes. The spectrum of the bis(β -phenylethyl)rhodamine was almost identical with that of diethylrhodamine.

"Rhodamine Dyes and Related Compounds. VI. Chloride and Amides of Sulforhodamine B," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1962), 32, 1489-92, the disclosure of which is totally incorporated herein by reference, discloses that sulforhodamine B (5 g., dried at 125°) and 3 g. PCl_5 heated in 50 milliliters CHCl_3 for 4 hours, then extd. with cold H_2O to remove excess PCl_6 , gave, after concentration of the dried organic

layer and treatment of the residue with much cold petroleum ether, the dark red p-sulfonyl chloride, $C_{27}H_{29}O_6N_2S_2Cl$, which slowly forms the original compound on contact with H_2O . With NH_3 in $CHCl_3$ it gave the corresponding p-sulfonamide, 81%, red-violet powder, sol. in EtOH or
5 AcOH; similarly was prepared the p-sulfonanilide, brown-violet solid. These have absorption spectra similar to the original compound but with less intense absorption. The p-sulfonyl chloride has a more intense absorption than the amides.

"Rhodamine Dyes and Related Compounds. V. α -Pyridylrhodamine," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1962), 32, 1485-9, the disclosure of which is totally incorporated herein by reference, discloses a process wherein heating 3,6-dichlorofluorane with 2-aminopyridine in the presence of $ZnCl_2$ for 3 hours at $160-80^\circ$ gave, after extraction with hot H_2O and EtOH and crystallization of the residue from
15 aqueous Me_2CO , 3-chloro-6- α -pyridylaminofluorane-HCl, m. $280-2^\circ$; free base, m. $185-7^\circ$. This heated with 2-aminopyridine and $ZnCl_2$ at $250-60^\circ$ for 6 hours, then precipitated from hot EtOH-HCl with H_2O , gave red N,N'-bis(α -pyridyl)rhodamine-HCl, m. $238-40^\circ$, also formed directly from dichlorofluorane and excess aminopyridine at $250-60^\circ$. Similarly,
20 3-chloro-6-anilino-fluorane gave red-violet N-phenyl-N'- α -pyridylrhodamine-HCl, m. $225-30^\circ$. All these were converted to N,N'-diphenylrhodamine by heating with $PhNH_2$ and $ZnCl_2$ for 3 hours at $180-200^\circ$. The absorption spectra of the products are shown; dipyrldylrhodamine has a more intense color than other
25 members of the group.

"Rhodamine Dyes and Related Compounds. IV. Aryl- and Benzylrhodamines," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1962), 32, 1480-5,

the disclosure of which is totally incorporated herein by reference, discloses a process wherein heating fluorescein chloride with ArNH_2 in the presence of $\text{ZnCl}_2\text{-ZnO}$ for 4 to 5 hours at $210\text{-}20^\circ$ gave, after leaching with hot dil. HCl, soln. of the residue in hot PhNH_2 , and pptn. with dil. HCl, the following N,N'-diarylrhodamines which were isolated as HCl salts: Ph, m. $255\text{-}60^\circ$; o-meC₆H₄, m. $205\text{-}10^\circ$; m-meC₆H₄, m. $195\text{-}200^\circ$; p-meC₆H₄, m. $255\text{-}60^\circ$. PhCH_2NH_2 similarly gave N,N'-dibenzylrhodamine, m. $160\text{-}5^\circ$; HCl salt decomp. $160\text{-}5^\circ$; di-HCl salt decomp. 210° . PhCH_2NH_2 and 3-chloro-6-anilino-2-fluoranthene gave 90-5% N-phenyl-N'-benzylrhodamine isolated as the HCl salt, m. $200\text{-}10^\circ$. The absorption spectra of these rhodamines are shown. Dibenzylrhodamine fluoresces strongly in solution, while the phenyl benzyl analog has a weak fluorescence. The benzyl groups cause a bathochromic shift of the absorption band in the substituted rhodamines; the diarylrhodamines form blue-violet solutions unlike the orange-yellow produced by unsubstituted rhodamine. The di-HCl salt of dibenzylrhodamine loses one HCl in soln. as shown by behavior in EtOH.

"Rhodamine Dyes and Related Compounds. III. Reaction of m-aminophenol With Phthalic Anhydride in Hot Sulfuric Acid," I. S. Ioffe et al., *Zh. Obsch. Khim.* (1962), 32, 1477-80, the disclosure of which is totally incorporated herein by reference, discloses that heating 25 g. of m-H₂NC₆H₄OH with 20 g. o-C₆H₄(CO)₂O in 100 milliliters concentrated H₂SO₄ at $160\text{-}200^\circ$ for 2-8 hours was used to examine the effects of conditions of condensation on the reaction products. Rhodamine formation began at 170° and reached a max. (20%) in 2 hours at 190° . Rhodol was a constant byproduct as a result of partial deamination of rhodamine. The deamination is promoted by longer reaction time and

higher temperatures. These factors also promoted the formation of a dark, amorphous material. o-Hydroxysulfanilic acid was formed in the reaction in up to 32% yield at 160° in 2 hours; more drastic conditions lowered its yield rapidly. Prior to the appearance of substantial amounts of rhodamine in the mixture, sulfonation of m-H₂NC₆H₄OH takes place, and the resulting compound appears to be the intermediate which reacts, with this compound forming rhodamine by displacement of the sulfonic acid group. This was confirmed by reaction of o-C₆H₄(CO)₂O with o-hydroxysulfanilic acid under the conditions shown above. m-Aminosalicylic acid also yields the same products in a mixture similar to that formed by m-H₂NC₆H₄OH.

"Rhodamine Dyes and Related Compounds. XVIII. N,N'-Dialkylrhodamines with Long Chain Hydrocarbon Radicals," I. S. Ioffe et al., *Zh. Organ. Khim.* (1970), 6(2), 369-71, the disclosure of which is totally incorporated herein by reference, discloses a process wherein the condensation of I (X=Cl) with RNH₂ (R=C₆H₁₃, C₈H₁₇, C₁₆H₃₃, or C₁₈H₃₇) gave the title dyes (I, X=NHR) (II). The presence of alkyl groups in II did not change their color in comparison with II (R=H); all II absorbed strongly at 523-6 nm. However, long alkyl chains altered the hydrophobic properties of II as shown by the change of their partition coefficients in oil-alc. or kerosine-alc. systems with the length of R chain.

"Rhodamine Dyes and Related Compounds. XIX. Mutual Transformations of Colorless and Colored Forms of N,N'-Substituted Rhodamine," I. S. Ioffe et al., *Zh. Organ. Khim.* (1972), 8(8), 1726-9, the disclosure of which is totally incorporated herein by reference, discloses that substituted rhodamines give colored solutions in polar and colorless solutions in nonpolar solvents. The solvent polarity at which the colorless

lactone form is converted to the quinoid, internal salt form depends on the number and structure of alkyl, aryl, or H substituents. Absorption spectra of N,N'-diethylrhodamine in water-dioxane mixtures show how the light absorption increases when the solvent polarity (i.e., water amount in the mixture) is increased.

"Synthesis of N-Substituted Flaveosines, Acridine Analogs of Rhodamine Dyes," I. S. Ioffe et al., *Zh. Org. Khim.* (1966), 2(9), 1721, the disclosure of which is totally incorporated herein by reference, discloses that o-(3,6-chloro-9-acridinyl)benzoic acid heated with BuNH₂ or Bu₂NH readily gave the hydrochlorides.

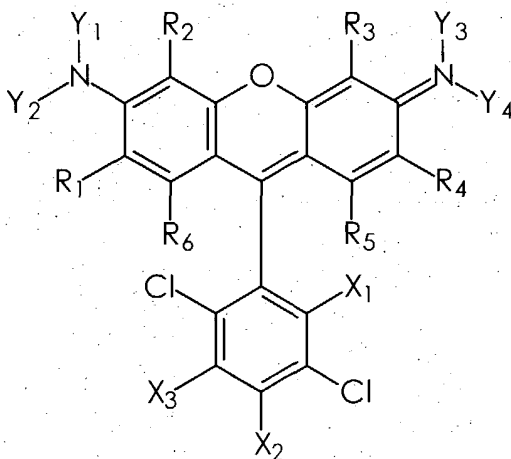
"Rhodamine Dyes and Related Compounds. XVII. Acridine Analogs of Rhodamine and Fluorescein," I. S. Ioffe et al., *Zh. Organ. Khim.* (1966), 2(5), 927-31, the disclosure of which is totally incorporated herein by reference, discloses absorption spectra for flaveosin, fluorescein, azafluorescein, their Et esters and diacetyl derivatives. Replacement of the xanthene structure by the acridine group changed the spectra of such dyes. Azafluorescein heated with PCl₅ at 95-100° gave o-(3,6-dichloro-9-acridinyl)-benzoic acid, decomp. >300°; its uv spectrum was similar to that of unsubstituted acridinylbenzoic acid. One of the flaveosin compounds heated with 25% H₂SO₄ in a sealed tube 10 hours at 200-20° gave azafluorescein, decomp. >380°; heated with EtOH-H₂SO₄ it gave one of the flaveosins, decomp. >300° Ac₂O-H₂SO₄ gave in 1 hour one of the flaveosins, decomp. 206°. The compound formed by treatment of 3,6-dichlorofluorane with NH₃ was prepared. Its uv spectrum is given.

"New Lipophilic Rhodamines and Their Application to Optical Potassium Sensing," T. Werner et al., *Journal of Fluorescence*,

Vol. 2, No. 3, pp. 93-98 (1992), the disclosure of which is totally incorporated herein by reference, discloses the synthesis of new lipophilic fluorescent rhodamines directly from 3,6-dichlorofluoresceins and the respective long-chain amines with excellent solubility in lipids and lipophilic membranes. Spectrophotometric and luminescent properties of the dyes are reported and discussed with respect to their application in new optical ion sensors. One rhodamine was applied in a poly(vinyl chloride)-based sensor membrane for continuous and sensitive optical determination of potassium ion, using valinomycin as the neutral ion carrier.

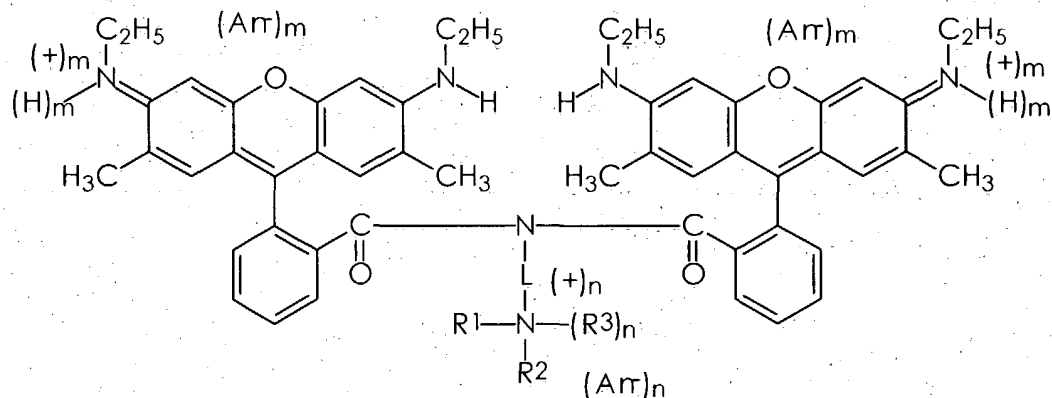
U.S. Patent 1,991,482 (Allemann), the disclosure of which is totally incorporated herein by reference, discloses a process of producing rhodamine dyes which comprises condensing a halogenated primary amine of the benzene series with fluorescein dichloride and sulfonating the condensed product.

U.S. Patent 5,847,162 (Lee et al.), the disclosure of which is totally incorporated herein by reference, discloses a class of 4,7-dichlororhodamine compounds useful as fluorescent dyes having the structure



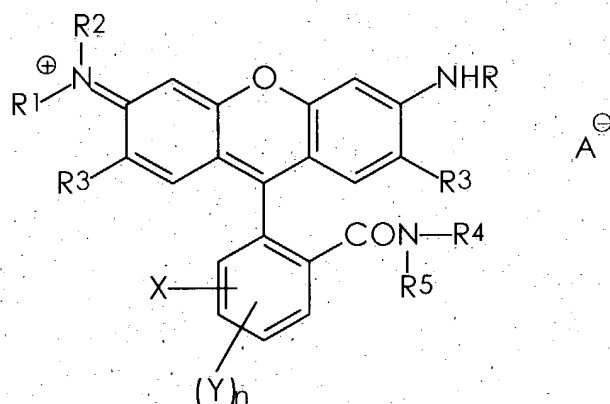
wherein R₁-R₆ are hydrogen, fluorine, chlorine, lower alkyl lower alkene, lower alkyne, sulfonate, sulfone, amino, amido, nitrile, lower alkoxy, lining group, or combinations thereof or, when taken together, R₁ and R₆ is benzo, or, when taken together, R₄ and R₅ is benzo; Y₁-Y₄ are hydrogen or lower alkyl or, when taken together, Y₁ and R₂ is propano and Y₂ and R₁ is propano, or, when taken together, Y₃ and R₃ is propano and Y₄ and R₄ is propano; and X₁-X₃ taken separately are selected from the group consisting of hydrogen, chlorine, fluorine, lower alkyl carboxylate, sulfonic acid, -CH₂OH, and linking group. In another aspect, the invention includes reagents labeled with the 4,7-dichlororhodamine dye compounds, including deoxynucleotides, dideoxynucleotides, and polynucleotides. In an additional aspect, the invention includes methods utilizing such dye compounds and reagents including dideoxy polynucleotide sequencing and fragment analysis methods.

U.S. Patent 4,935,059 (Mayer et al.), the disclosure of which is totally incorporated herein by reference, discloses basic rhodamine dyes suitable for use in recording fluids for the ink jet process and for coloring paper stock having the formula



- where L is C₂-C₁₀-alkylene, R¹, R², and R³ are each independently of the others hydrogen, substituted or unsubstituted C₁-C₁₀-alkyl or C₅-C₇-cycloalkyl or R¹ and R² together with the nitrogen atom linking them together are a hetero cyclic radical, A⁻ is one equivalent of an anion and m and n are each independently of the other 0 or 1.

U.S. Patent 4,647,675 (Mayer et al.), the disclosure of which is totally incorporated herein by reference, discloses compounds of the general formula



- 10 where A⁻ is an anion, R is hydrogen or unsubstituted or substituted alkyl or cycloalkyl, R¹ and R² independently of one another are each hydrogen or unsubstituted or substituted alkyl or cycloalkyl, or one of the radicals may furthermore be aryl, or R¹ and R², together with the nitrogen atom, form a saturated heterocyclic structure, the radicals R³
- 15 independently of one another are each hydrogen or C₁-C₄-alkyl, R⁴ and R⁵ independently of one another are each unsubstituted or substituted alkyl or cycloalkyl, or one of the radicals may furthermore be hydrogen, aryl or hetaryl, R⁴ and R⁵, together with the nitrogen atom, form a saturated heterocyclic structure, n is 1, 2 or 3, X is hydrogen,

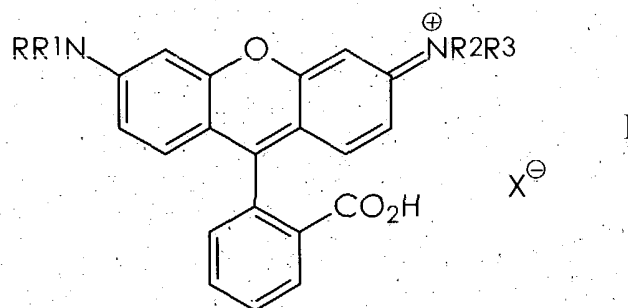
chlorine, bromine, C₁-C₄-alkyl, C₁-C₄-alkoxy or nitro and Y is hydrogen or chlorine, are particularly useful for dyeing paper stocks.

U.S. Patent 1,981,515 (Kyrides), the disclosure of which is totally incorporated herein by reference, discloses intermediates for rhodamine dyestuffs.

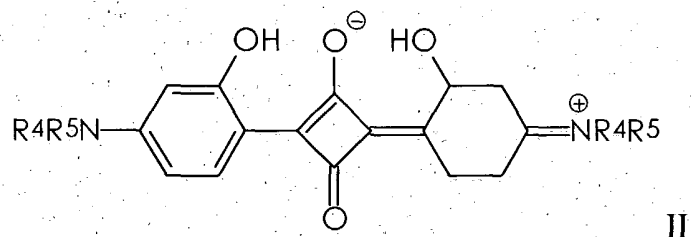
U.S. Patent 1,981,516 (Kyrides), the disclosure of which is totally incorporated herein by reference, discloses intermediates for secondary alkylated rhodamine dyes.

British Patent Publication GB 421 737, the disclosure of which is totally incorporated herein by reference, discloses dyes of the rhodamine series which are prepared by condensing naphthalene-2:3-dicarboxylic acid with a m-aminophenol in which the nitrogen group is substituted by one or two alkyl groups, the products, if desired, being sulphonated. The unsulphonated products may be used as lake colouring matters whilst the sulphonated dyes are acid wool dyes. In examples, (1) naphthalene-2:3-dicarboxylic acid is condensed with diethyl-m-aminophenol in the presence of zinc chloride giving a product which dyes tannin-mordanted cotton in the same shade as Rhodamine B and a sulphonated product which dyes wool bluish-red shades; (2) monoethyl-m-aminophenol is used instead of the diethyl-m-aminophenol in example (1), yielding a dye, which when sulphonated dyes wool red-orange shades; (3) 2-ethylamino-p-cresol replaces the diethyl-m-aminophenol in example (1), yielding a dye dyeing and printing tannin-mordanted cotton in shades similar to Rhodamine 69BS and when sulphonated dyeing wool red.

Japanese Patent Publication JP 61221265, the disclosure of which is totally incorporated herein by reference, discloses rhodamine compounds of formula I



- 5 wherein R₁, R₃ are each lower alkyl; R₂ is lower alkyl, 10C or higher long-chain alkyl; R₄ is 10C or higher long-chain alkyl; X⁻ is an anion, or squarylium compounds of formula II



- wherein R₂ is 10C or higher long-chain alkyl. Example: 3,6-(N,N'-diethyl-N,N'-dioctadecyl) diamino-9-(2-carboxyphenyl) xanthilium perchlorate.
- 10 Use: materials for molecular electronics, which are suitable for use as materials for photoelectric converter, optical memory, etc.
- Preparation: 2-(4-N,N'-diethylamino-2-hydroxybenzoyl)-benzoic acid, which is a condensate between N-ethyl-N-octadecyl-m-hydroxyaniline
- 15 and phthalic anhydride, is reacted with N-ethyl-N-octadecyl-m-hydroxyaniline to obtain the compound of formula I.
- 3-HOC₆H₄N(Et)(CH₂)₁₇Me and phthalic anhydride were heated at 150°

for 4 hours, treated with aqueous NH_3 , and the amorphous intermediate mixed with aqueous HClO_4 forming a compound of formula I ($\text{R}=\text{R}_2=\text{Et}$; $\text{R}_1=\text{R}_3=\text{C}_{18}\text{H}_{37}$; $\text{X}=\text{ClO}_4$), having λ_{max} (MeOH) 550 nm.

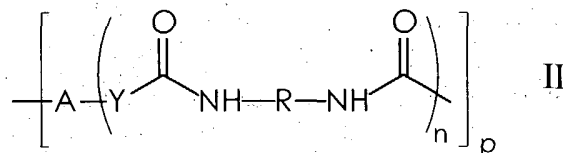
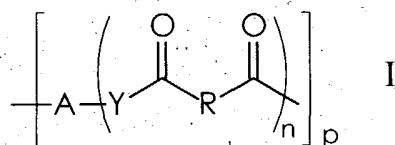
U.S. Patent 5,084,099 (Jaeger et al.), the disclosure of which
5 is totally incorporated herein by reference, discloses modified phase change ink compatible colorants which comprise a phase change ink soluble complex of (a) a tertiary alkyl primary amine and (b) dye chromophores, i.e., materials that absorb light in the visible wavelength region to produce color having at least one pendant acid functional
10 group in the free acid form (not the salt of that acid). These modified colorants are extremely useful in producing phase change inks when combined with a phase change ink carrier, even though the unmodified dye chromophores have limited solubility in the phase change ink carrier. Thin films of uniform thickness of the subject phase
15 change ink compositions which employ the modified phase change ink colorants exhibit a high degree of lightness and chroma. The primary amine-dye chromophore complexes are soluble in the phase change ink carrier and exhibit excellent thermal stability.

U.S. Patent 5,507,864 (Jaeger et al.), the disclosure of which
20 is totally incorporated herein by reference, discloses a phase change ink composition that includes a combination of different dye types such as an anthraquinone dye and a xanthene dye, which is most preferably a rhodamine dye. While each dye type is insufficiently soluble with respect to favored carrier compositions to preserve color saturation in
25 reduced ink quantity prints, the dye type combination permits increased dye loading and maintains print quality. In a preferred embodiment of the invention, a favored carrier composition is adjusted

to promote the colored form of a preferred rhodamine dye (C.I. Solvent Red 49) and mixed with a preferred anthraquinone dye (C.I. Solvent Red 172) whose concentration is kept below a critical level to prevent post printed blooming. The resulting preferred phase change ink compositions provide a magenta phase change ink with enhanced light fastness and color saturation, as well as good compatibility with preferred existing subtractive primary color phase change inks.

U.S. Patent 5,621,022 (Jaeger et al.), the disclosure of which is totally incorporated herein by reference, discloses a phase change ink composition wherein the ink composition utilizes polymeric dyes in combination with a selected phase change ink carrier composition.

U.S. Patent 5,747,554 (Sacripante et al.), the disclosure of which is totally incorporated herein by reference, discloses an ink composition comprising a polyesterified-dye (I) or polyurethane-dye (II) with a viscosity of from about 3 centipoise to about 20 centipoise at a temperature of from about 125°C to about 165°C and represented by the formulas



wherein A is an organic chromophore, Y is an oxyalkylene or poly(oxyalkylene), R is an arylene or alkylene, n represents the number of repeating segments, and is an integer of from about 2 to about 50,

and p represents the number of chains per chromophore and is an integer of from about 1 to about 6.

U.S. Patent 5,902,841 (Jaeger et al.), the disclosure of which is totally incorporated herein by reference, discloses a phase change ink composition wherein the ink composition utilizes colorant in combination with a selected phase change ink carrier composition containing at least one hydroxy-functional fatty amide compound.

European Patent Publication 0 565 798 (Shustack), the disclosure of which is totally incorporated herein by reference, discloses ultraviolet radiation-curable primary and secondary coating compositions for optical fibers. The primary coatings comprise a hydrocarbon polyol-based reactively terminated aliphatic urethane oligomer; a hydrocarbon monomer terminated with at least one end group capable of reacting with the terminus of the oligomer; and an optional photoinitiator. The secondary coatings comprise a polyester and/or polyether-based aliphatic urethane reactively terminated oligomer; a hydrocarbonaceous viscosity-adjusting component capable of reacting with the reactive terminus of (I); and an optional photoinitiator. Also disclosed are optical fibers coated with the secondary coating alone or with the primary and secondary coatings of the invention.

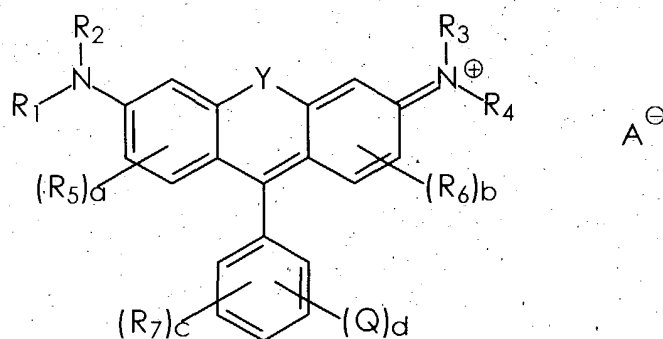
While known compositions and processes are suitable for their intended purposes, a need remains for new magenta colorant compositions. In addition, a need remains for magenta colorant compositions particularly suitable for use in phase change inks. Further, a need remains for magenta colorants with desirable thermal stability. Additionally, a need remains for magenta colorants that exhibit minimal

undesirable discoloration when exposed to elevated temperatures. There is also a need for magenta colorants that exhibit a desirable brilliance. In addition, there is a need for magenta colorants that exhibit a desirable hue. Further, there is a need for magenta colorants that are of desirable chroma. Additionally, there is a need for magenta colorants that have desirably high lightfastness characteristics. A need also remains for magenta colorants that have a desirably pleasing color. In addition, a need remains for magenta colorants that exhibit desirable solubility characteristics in phase change ink carrier compositions. Further, a need remains for magenta colorants that enable phase change inks to be jetted at temperatures of over 135°C while maintaining thermal stability. Additionally, a need remains for magenta colorants that enable phase change inks that generate images with low pile height. There is also a need for magenta colorants that enable phase change inks that generate images that approach lithographic thin image quality. In addition, there is a need for magenta colorants that exhibit oxidative stability. Further, there is a need for magenta colorants that do not precipitate from phase change ink carriers. Additionally, there is a need for magenta colorants that do not, when included in phase change inks, diffuse into adjacently printed inks of different colors. A need also remains for magenta colorants that do not leach from media such as phase change ink carriers into tape adhesives, paper, or the like. In addition, a need remains for magenta colorants that, when incorporated into phase change inks, do not lead to clogging of a phase change ink jet printhead. Further, there is a need for magenta colorants that enable phase change inks that generate images with sharp edges that remain

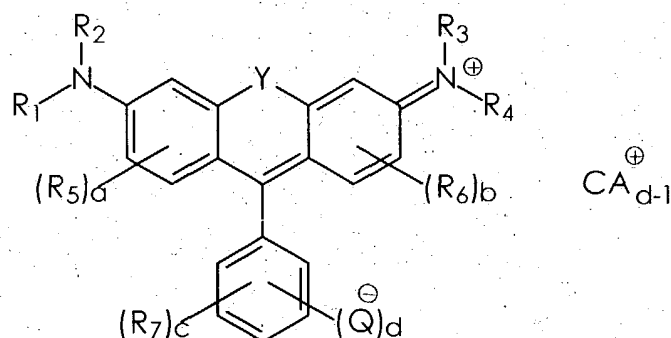
sharp over time. Additionally, there is a need for magenta colorants that enable phase change inks that generate images which retain their high image quality in warm climates. Further, there is a need for magenta colorants that enable phase change inks that generate images of desirably high optical density. Additionally, there is a need for magenta colorants that, because of their good solubility in phase change ink carriers, enable the generation of images of low pile height without the loss of desirably high optical density. A need also remains for magenta colorants that enable cost-effective inks.

SUMMARY

The present invention is directed to a phase change ink composition comprising a phase change ink carrier and a colorant compound of the formula

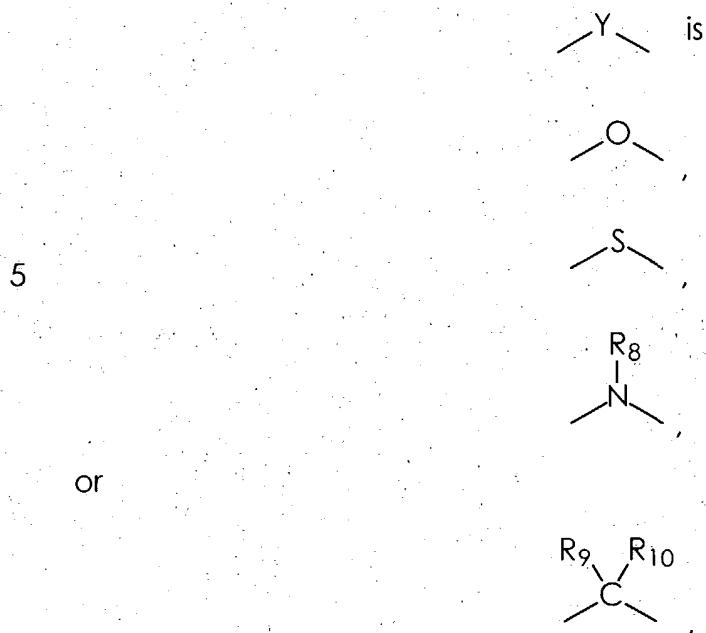


or



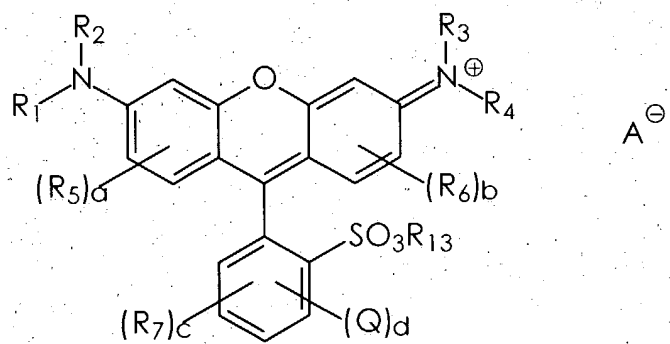
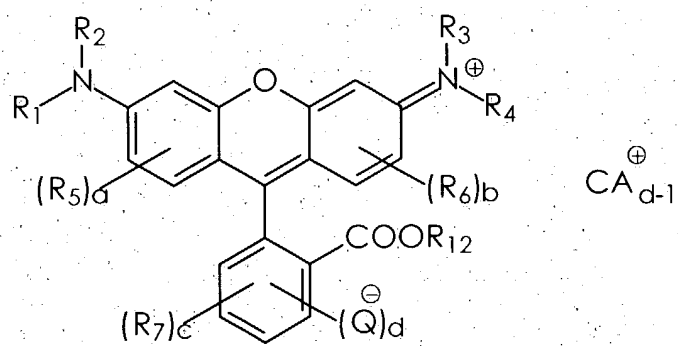
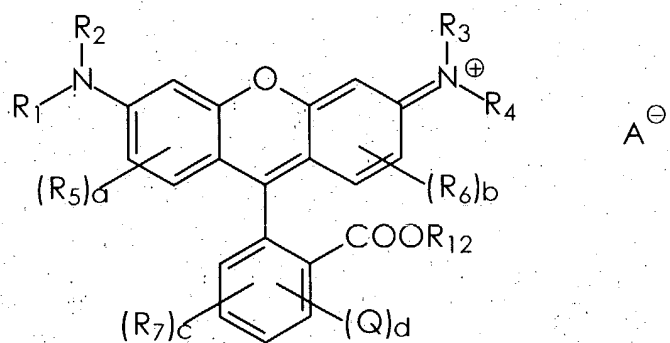
wherein R₁, R₂, R₃, and R₄ each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group, (iii) an aryl group, (iv) an arylalkyl group, or (v) an alkylaryl group, wherein R₁ and R₂ can be joined together to form a ring, wherein R₃ and R₄ can be joined together to form a ring, and wherein R₁, R₂, R₃, and R₄ can each be joined to a phenyl ring in the central structure, a and b each, independently of the others, is an integer which is 0, 1, 2, or 3, c is an integer which is 0, 1, 2, 3, or 4, each R₅, R₆, and R₇, independently of the others, is (i) an alkyl group, (ii) an aryl group, (iii) an arylalkyl group, (iv) an alkylaryl group, (v) a halogen atom, (vi) an ester group, (vii) an amide group, (viii) a sulfone group, (ix) an amine group or ammonium group, (x) a nitrile group, (xi) a nitro group, (xii) a hydroxy group, (xiii) a cyano group, (xiv) a pyridine or pyridinium group, (xv) an ether group, (xvi) an aldehyde group, (xvii) a ketone group, (xviii) a carbonyl group, (xix) a thiocarbonyl group, (xx) a sulfate group, (xxi) a sulfide group, (xxii) a sulfoxide group, (xxiii) a phosphine or phosphonium group, (xxiv) a phosphate group, (xxv) a mercapto group, (xxvi) a nitroso group, (xxvii) an acyl group, (xxviii) an acid anhydride group, (xxix) an azide group, (xxx) an azo group, (xxxi) a cyanato group, (xxxii) an isocyanato group, (xxxiii) a thiocyanato group, (xxxiv) an isothiocyanato group, (xxxv) a

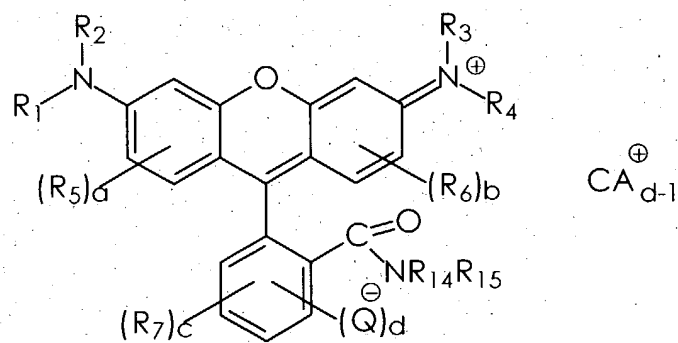
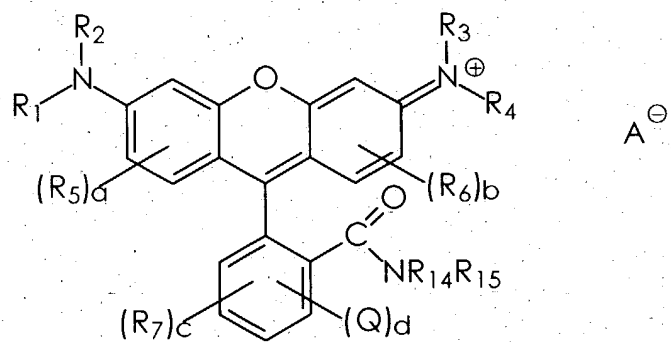
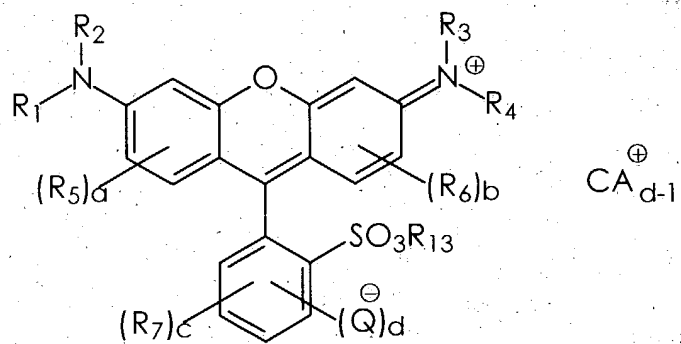
urethane group, or (xxxvi) a urea group, wherein R_5 , R_6 , and R_7 can each be joined to a phenyl ring in the central structure,

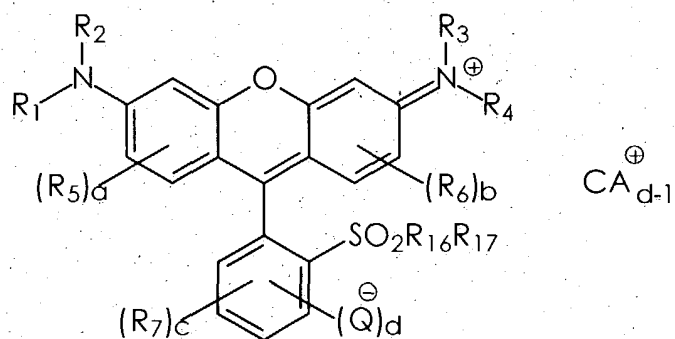
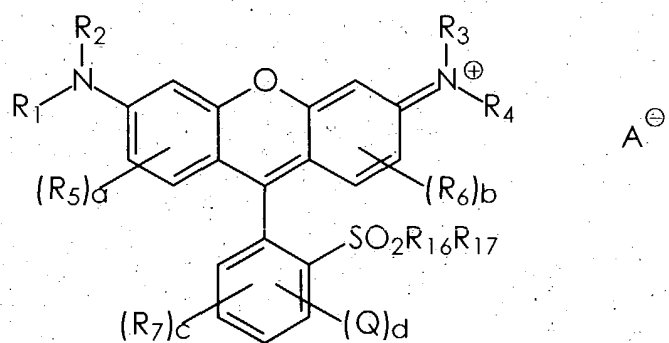


10 R_8 , R_9 , and R_{10} each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group, (iii) an aryl group, (iv) an arylalkyl group, or (v) an alkylaryl group, provided that the number of carbon atoms in $R_1+R_2+R_3+R_4+R_5+R_6+R_7+R_8+R_9+R_{10}$ is at least about 16, Q is a COOH group or a SO₃H group, Q⁻ is a COO⁻ group or a SO₃⁻ group, d is an integer which is 1, 2, 3, 4, or 5, A is an anion, and CA is either a

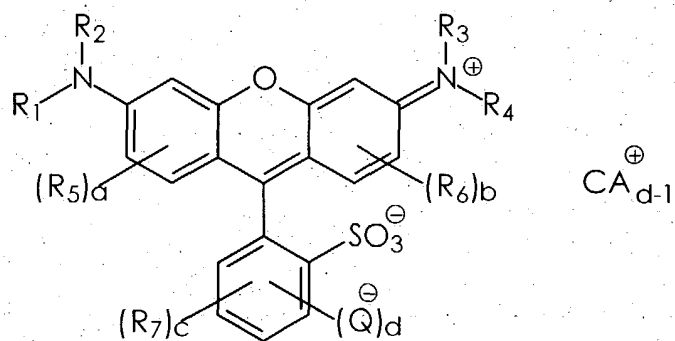
15 hydrogen atom or a cation associated with all but one of the Q⁻ groups, provided that when the colorant compound is of the formula







or

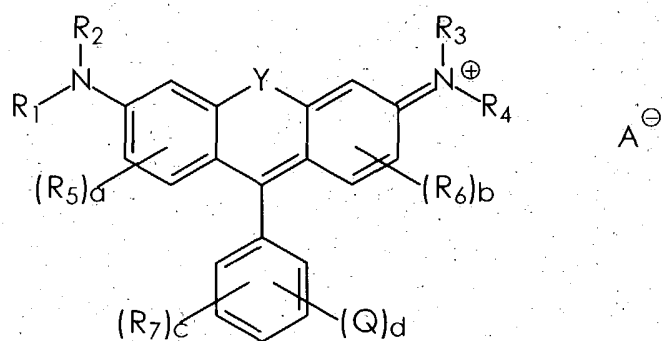


- 5 wherein R_{12} , R_{13} , R_{14} , R_{15} , R_{16} , and R_{17} each, independently of the other, is (i) an alkyl group, (ii) an aryl group, (iii) an arylalkyl group, or (iv) an alkylaryl group, (I) either (a) c is an integer which is 0, 1, 2, or 3, or (b) d

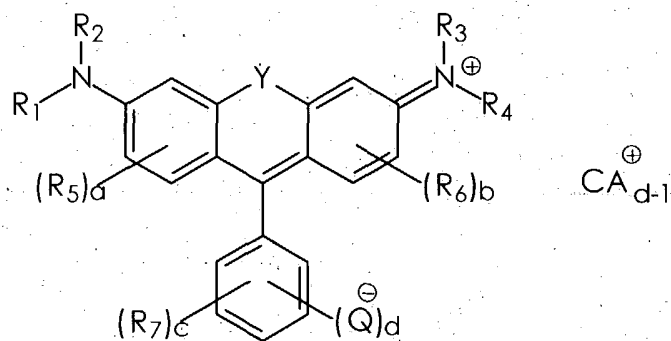
is an integer which is 1, 2, 3, or 4, and (II) either (a) three of R_1 , R_2 , R_3 , and R_4 are hydrogen atoms; (b) only one of R_1 , R_2 , R_3 , and R_4 is a hydrogen atom; (c) R_1 and R_2 are both hydrogen atoms; (d) R_3 and R_4 are both hydrogen atoms; or (e) R_1 and R_3 are both hydrogen atoms and R_2 and R_4 are each, independently of the other, either alkyl groups or arylalkyl groups.

DETAILED DESCRIPTION

The present invention is directed to phase change inks containing colorant compounds of the formula



or

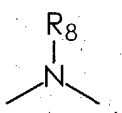
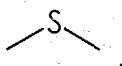
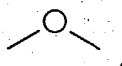
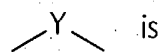


wherein R₁, R₂, R₃, and R₄ each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group (including linear, branched, saturated, unsaturated, cyclic, substituted, and unsubstituted alkyl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkyl group), in one embodiment with at least 1 carbon atom, in another embodiment with at least about 2 carbon atoms, in yet another embodiment with at least about 6 carbon atoms, in another embodiment with at least about 8 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iii) an aryl group (including unsubstituted and substituted aryl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the aryl group), in one embodiment with at least about 6 carbon atoms, in another embodiment with at least about 10 carbon atoms, and in yet another embodiment with at least about 14 carbon atoms, and in one embodiment with no more than about 26 carbon atoms, in another embodiment with no more than about 22 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iv) an arylalkyl group (including unsubstituted and substituted arylalkyl groups, wherein the alkyl portion of the arylalkyl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero

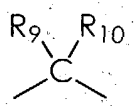
atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the arylalkyl group), in one embodiment with at least about 7 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as benzyl or the like, or (v) an alkylaryl group (including unsubstituted and substituted alkylaryl groups, wherein the alkyl portion of the alkylaryl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylaryl group), in one embodiment with at least about 7 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, wherein R₁ and R₂ can be joined together to form a ring, wherein R₃ and R₄ can be joined together to form a ring, and wherein R₁, R₂, R₃, and R₄ can each be joined to a phenyl ring in the central structure, a and b each, independently of the others, is an integer which is 0, 1, 2, or 3, c is an

integer which is 0, 1, 2, 3, or 4, each R_5 , R_6 , and R_7 , independently of the others, is (i) an alkyl group (including linear, branched, saturated, unsaturated, cyclic, substituted, and unsubstituted alkyl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkyl group), in one embodiment with at least 1 carbon atom, and in one embodiment with no more than about 50 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (ii) an aryl group (including unsubstituted and substituted aryl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the aryl group), in one embodiment with at least about 6 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iii) an arylalkyl group (including unsubstituted and substituted arylalkyl groups, wherein the alkyl portion of the arylalkyl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the arylalkyl group), in one embodiment with at least about 7 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no

more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as benzyl or the like, (iv) an alkylaryl group (including unsubstituted and substituted alkylaryl groups, wherein the alkyl portion of the alkylaryl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylaryl group), in one embodiment with at least about 7 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, (v) a halogen atom, such as fluorine, chlorine, bromine, iodine, or the like, (vi) an ester group, (vii) an amide group, (viii) a sulfone group, (ix) an amine group or ammonium group, (x) a nitrile group, (xi) a nitro group, (xii) a hydroxy group, (xiii) a cyano group, (xiv) a pyridine or pyridinium group, (xv) an ether group, (xvi) an aldehyde group, (xvii) a ketone group, (xviii) a carbonyl group, (xix) a thiocarbonyl group, (xx) a sulfate group, (xxi) a sulfide group, (xxii) a sulfoxide group, (xxiii) a phosphine or phosphonium group, (xxiv) a phosphate group, (xxv) a mercapto group, (xxvi) a nitroso group, (xxvii) an acyl group, (xxviii) an acid anhydride group, (xxix) an azide group, (xxx) an azo group, (xxxi) a cyanato group, (xxxii) an isocyanato group, (xxxiii) a thiocyanato group, (xxxiv) an isothiocyanato group, (xxxv) a urethane group, or (xxxvi) a urea group, wherein R₅, R₆, and R₇ can each be joined to a phenyl ring in the central structure,



5 or

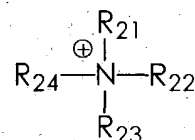


R_8 , R_9 , and R_{10} each, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group (including linear, branched, saturated, unsaturated, cyclic, substituted, and unsubstituted alkyl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkyl group), in one embodiment with at least 1 carbon atom, in another embodiment with at least about 2 carbon atoms, in yet another embodiment with at least about 6 carbon atoms, in another embodiment with at least about 8 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iii) an aryl group (including unsubstituted and substituted aryl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon,

phosphorus, and the like either may or may not be present in the aryl group), in one embodiment with at least about 6 carbon atoms, in another embodiment with at least about 10 carbon atoms, and in yet another embodiment with at least about 14 carbon atoms, and in one
5 embodiment with no more than about 26 carbon atoms, in another embodiment with no more than about 22 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iv) an arylalkyl group (including unsubstituted and substituted arylalkyl
10 groups, wherein the alkyl portion of the arylalkyl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the arylalkyl group), in one embodiment with at
15 least about 7 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more
20 than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as benzyl or the like, or (v) an alkylaryl group (including unsubstituted and substituted alkylaryl groups, wherein the alkyl portion of the alkylaryl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such
25 as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylaryl group), in one embodiment with at least about 7

carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, provided that the number of carbon atoms in $R_1+R_2+R_3+R_4+R_5+R_6+R_7+R_8+R_9+R_{10}$ is in one embodiment at least about 16, in another embodiment at least about 18, in yet another embodiment at least about 20, in still another embodiment at least about 22, in another embodiment at least about 24, in yet another embodiment at least about 26, in still another embodiment at least about 28, in another embodiment at least about 30, in yet another embodiment at least about 32, in still another embodiment at least about 34, in another embodiment at least about 36, in yet another embodiment at least about 38, in still another embodiment at least about 40, in another embodiment at least about 42, in yet another embodiment at least about 44, in still another embodiment at least about 46, in another embodiment at least about 48, in yet another embodiment at least about 50, in still another embodiment at least about 52, in another embodiment at least about 54, in yet another embodiment at least about 56, in still another embodiment at least about 58, in another embodiment at least about 60, in yet another embodiment at least about 62, in still another embodiment at least about 64, in another embodiment at least about 66, in yet another embodiment at least about 68, in still another embodiment at least about 70, and in another embodiment at least about 72, each Q,

independently of the others, is a COOH group or a SO₃H group, each Q⁻, independently of the others, is a COO⁻ group or a SO₃⁻ group, d is an integer which is 1, 2, 3, 4, or 5, A is an anion, with examples of suitable anions including (but not being limited to) Cl⁻, Br⁻, I⁻, HSO₄⁻, HSO₃⁻, SO₄²⁻,
 5 SO₃²⁻, CH₃SO₃⁻, CH₃C₆H₄SO₃⁻, NO₃⁻, HCOO⁻, CH₃COO⁻, H₂PO₄⁻, HPO₄²⁻, SCN⁻, BF₄⁻, ClO₄⁻, SSO₃⁻, PF₆⁻, SbCl₆⁻, or the like, as well as mixtures thereof, and CA is either a hydrogen atom or a cation associated with all but one of the Q⁻ groups, with examples of suitable cations including (but not being limited to) alkali metal cations, such as Li⁺, Na⁺, K⁺, Rb⁺, and
 10 Cs⁺, nonpolymeric or monomeric ammonium and quaternary amine cations, including those of the general formula

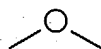


wherein each of R₂₁, R₂₂, R₂₃, and R₂₄, independently of the others, is (i) a hydrogen atom, (ii) an alkyl group (including linear, branched, saturated, unsaturated, cyclic, substituted, and unsubstituted alkyl
 15 groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkyl group), in one embodiment with at least 1 carbon atom, in another embodiment with at least about 2 carbon atoms, in yet another embodiment with at least about 6 carbon atoms, in another
 20 embodiment with at least about 8 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet
 25 another embodiment with no more than about 20 carbon atoms,

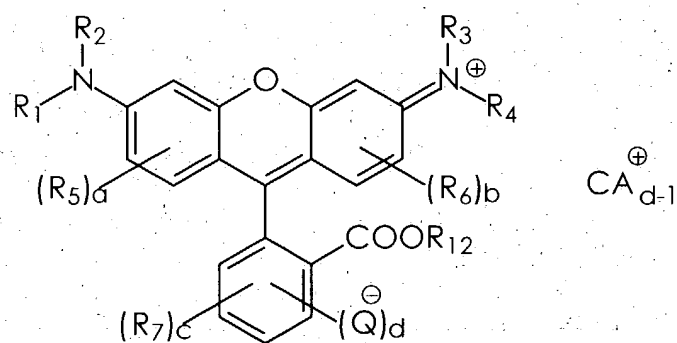
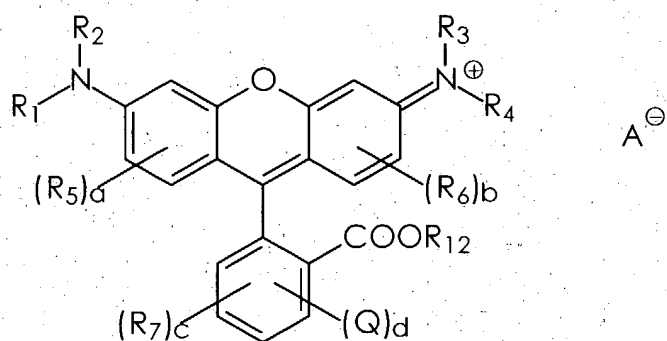
although the number of carbon atoms can be outside of these ranges,
(iii) an aryl group (including unsubstituted and substituted aryl groups,
and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon,
phosphorus, and the like either may or may not be present in the aryl
5 group), in one embodiment with at least about 6 carbon atoms, in
another embodiment with at least about 10 carbon atoms, and in yet
another embodiment with at least about 14 carbon atoms, and in one
embodiment with no more than about 26 carbon atoms, in another
embodiment with no more than about 22 carbon atoms, and in yet
10 another embodiment with no more than about 18 carbon atoms,
although the number of carbon atoms can be outside of these ranges,
(iv) an arylalkyl group (including unsubstituted and substituted arylalkyl
groups, wherein the alkyl portion of the arylalkyl group can be linear,
branched, saturated, unsaturated, and/or cyclic, and wherein hetero
15 atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like
either may or may not be present in either or both of the alkyl portion
and the aryl portion of the arylalkyl group), in one embodiment with at
least about 7 carbon atoms, in another embodiment with at least
about 12 carbon atoms, and in yet another embodiment with at least
20 about 18 carbon atoms, and in one embodiment with no more than
about 55 carbon atoms, in another embodiment with no more than
about 30 carbon atoms, and in yet another embodiment with no more
than about 20 carbon atoms, although the number of carbon atoms
can be outside of these ranges, such as benzyl or the like, or (v) an
25 alkylaryl group (including unsubstituted and substituted alkylaryl groups,
wherein the alkyl portion of the alkylaryl group can be linear, branched,
saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such

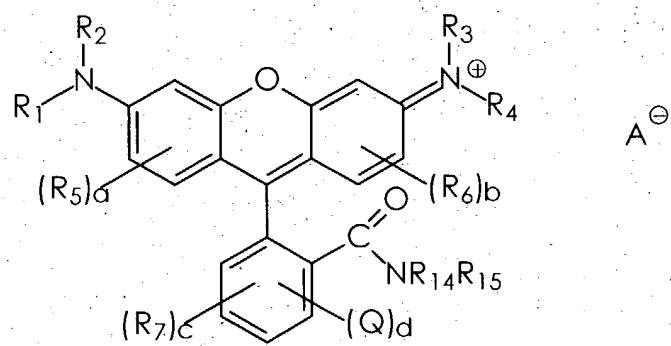
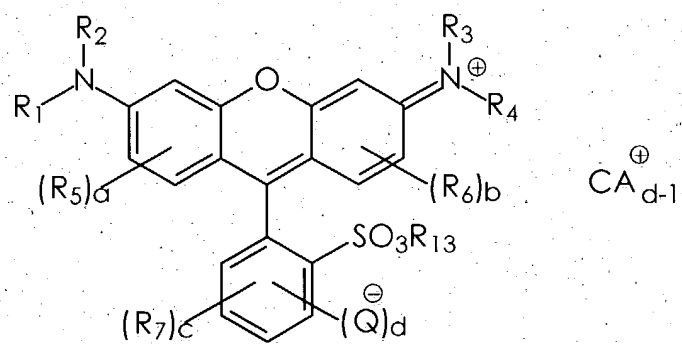
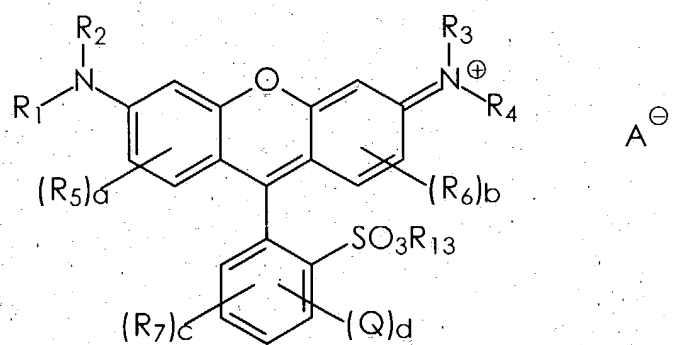
as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylaryl group), in one embodiment with at least about 7 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, wherein one or more of R₂₁, R₂₂, R₂₃, and R₂₄ can be joined together to form a ring, and wherein the substituents on the substituted alkyl, aryl, arylalkyl, and alkylaryl groups can be (but are not limited to) hydroxy groups, halogen atoms, amine groups, imine groups, ammonium groups, cyano groups, pyridine groups, pyridinium groups, ether groups, aldehyde groups, ketone groups, ester groups, amide groups, carbonyl groups, thiocarbonyl groups, sulfate groups, sulfonate groups, sulfonic acid groups, sulfide groups, sulfoxide groups, phosphine groups, phosphonium groups, phosphate groups, nitrile groups, mercapto groups, nitro groups, nitroso groups, sulfone groups, acyl groups, acid anhydride groups, azide groups, azo groups, cyanato groups, isocyanato groups, thiocyanato groups, isothiocyanato groups, carboxylate groups, carboxylic acid groups, urethane groups, urea groups, mixtures thereof, and the like, wherein two or more substituents can be joined together to form a ring, oligomeric and polymeric cations, such as cationic polymers or oligomers, and the like, as well as mixtures thereof.

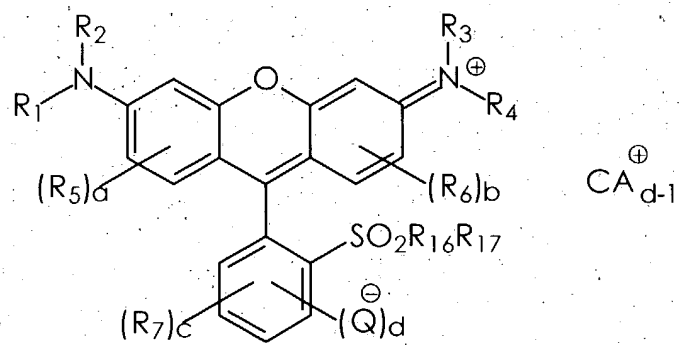
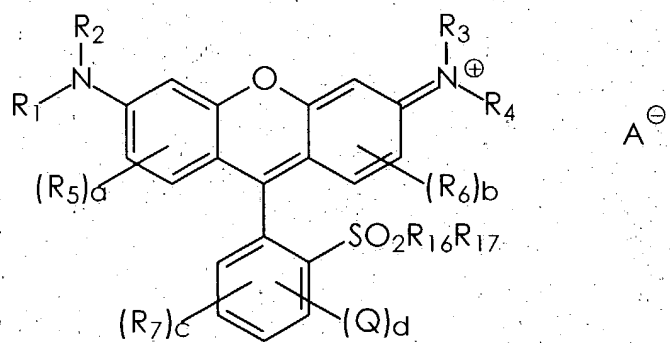
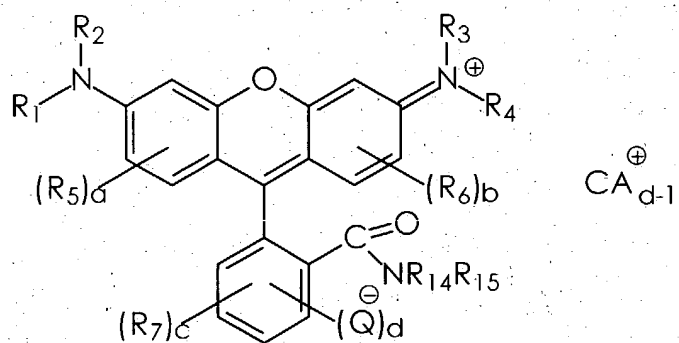
In situations wherein



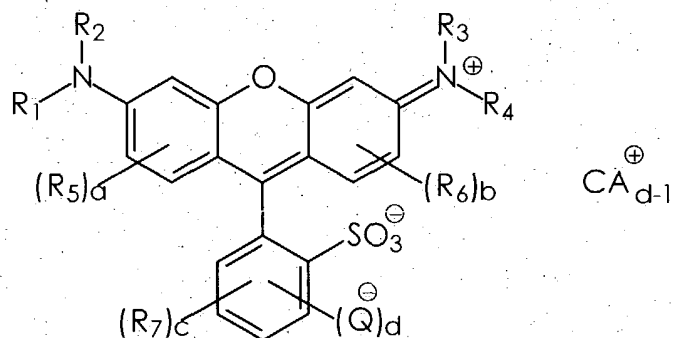
- and either (i) one of the R_7 groups is in the ortho position and is either an ester based on a carboxylic acid, an ester based on a sulfonic acid, an amide based on a carboxylic acid, or an amide based on a sulfonic acid, or (ii) one of the Q groups is a sulfonate salt, i.e., when the colorant compound is of the formula







or



wherein R_{12} , R_{13} , R_{14} , R_{15} , R_{16} , and R_{17} each, independently of the other, is

(i) an alkyl group (including linear, branched, saturated, unsaturated, cyclic, substituted, and unsubstituted alkyl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkyl group), in one embodiment with at least 1 carbon atom, and in one embodiment with no more than about 50 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (ii) an aryl group (including unsubstituted and substituted aryl groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the aryl group), in one embodiment with at least about 6 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges,

(iii) an arylalkyl group (including unsubstituted and substituted arylalkyl

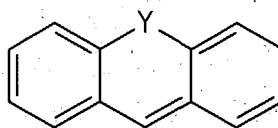
groups, wherein the alkyl portion of the arylalkyl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the arylalkyl group), in one embodiment with at least about 7 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as benzyl or the like, or (iv) an alkylaryl group (including unsubstituted and substituted alkylaryl groups, wherein the alkyl portion of the alkylaryl group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylaryl group), in one embodiment with at least about 7 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, wherein the substituents on the substituted alkyl, aryl, arylalkyl, and alkylaryl groups can be (but are not limited to) hydroxy groups, halogen atoms, amine groups, imine groups, ammonium groups, cyano groups, pyridine groups, pyridinium groups, ether groups, aldehyde groups, ketone groups, ester groups, amide groups, carbonyl groups, thiocarbonyl groups, sulfate groups, sulfonate groups, sulfonic acid groups, sulfide

groups, sulfoxide groups, phosphine groups, phosphonium groups, phosphate groups, nitrile groups, mercapto groups, nitro groups, nitroso groups, sulfone groups, acyl groups, acid anhydride groups, azide groups, azo groups, cyanato groups, isocyanato groups, thiocyanato groups, isothiocyanato groups, carboxylate groups, carboxylic acid groups, urethane groups, urea groups, mixtures thereof, and the like, wherein two or more substituents can be joined together to form a ring, (I) either (a) c is an integer which is 0, 1, 2, or 3, or (b) d is an integer which is 1, 2, 3, or 4, and (II) either (a) three of R₁, R₂, R₃, and R₄ are hydrogen atoms; (b) only one of R₁, R₂, R₃, and R₄ is a hydrogen atom; (c) R₁ and R₂ are both hydrogen atoms; (d) R₃ and R₄ are both hydrogen atoms; or (e) R₁ and R₃ are both hydrogen atoms and R₂ and R₄ are each, independently of the other, either alkyl groups or arylalkyl groups.

In one embodiment, the number of carbon atoms in R₁+R₂+R₃+R₄ is at least about 16, in another embodiment at least about 18, in yet another embodiment at least about 20, in still another embodiment at least about 22, in another embodiment at least about 24, in yet another embodiment at least about 26, in still another embodiment at least about 28, in another embodiment at least about 30, in yet another embodiment at least about 32, in still another embodiment at least about 34, in another embodiment at least about 36, in yet another embodiment at least about 38, in still another embodiment at least about 40, in another embodiment at least about 42, in yet another embodiment at least about 44, in still another embodiment at least about 46, in another embodiment at least about 48, in yet another embodiment at least about 50, in still another

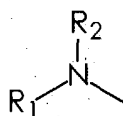
embodiment at least about 52, in another embodiment at least about 54, in yet another embodiment at least about 56, in still another embodiment at least about 58, in another embodiment at least about 60, in yet another embodiment at least about 62, in still another embodiment at least about 64, in another embodiment at least about 66, in yet another embodiment at least about 68, in still another embodiment at least about 70, and in another embodiment at least about 72.

Since hetero atoms can be included in the alkyl, aryl, arylalkyl, and alkylaryl groups, and since the groups can be substituted, it is to be understood that R_1 , R_2 , R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , and R_{10} can also be groups such as alkoxy, polyalkyleneoxy, aryloxy, polyaryleneoxy, arylalkyloxy, polyarylalkyleneoxy, alkylaryloxy, or polyalkylaryleneoxy groups, provided that the oxygen atom in such a group is not directly bonded to a nitrogen, oxygen, or sulfur atom in the



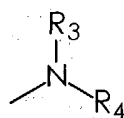
central structure.

Examples of situations wherein one of the R_{1-4} groups is a cycloalkyl is when

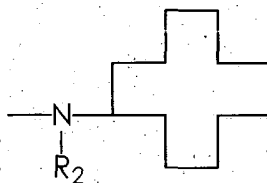


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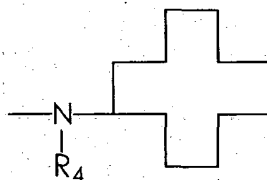
or



is

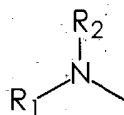


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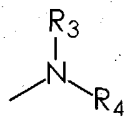


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Examples of situations wherein the R_{1-4} groups are joined together to form a ring are when

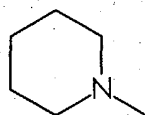


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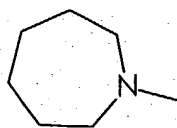


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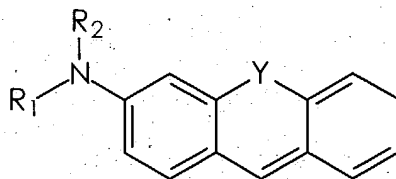
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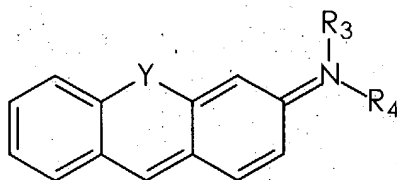
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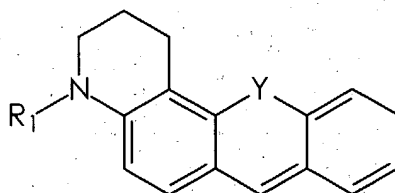
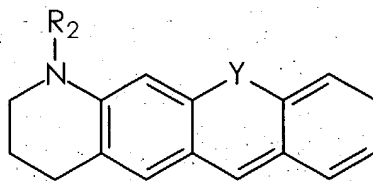
Examples of situations wherein one of the R_{1-4} groups is joined to a phenyl ring in the central structure is when

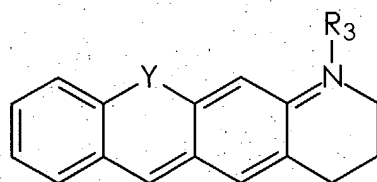


5 or

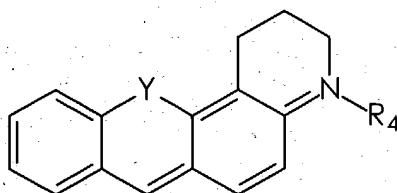


is

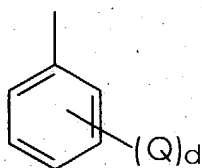




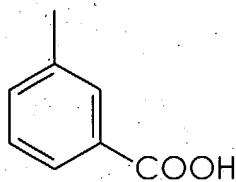
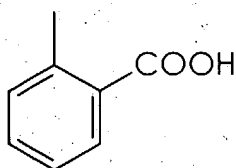
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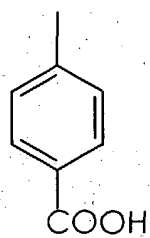
Compounds of the present invention include
 5 monocarboxylic acids and monocarboxylates, wherein



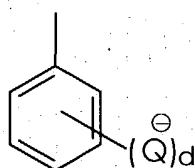
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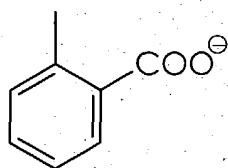
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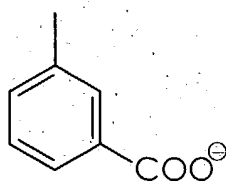
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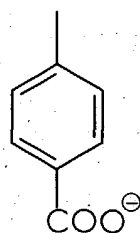
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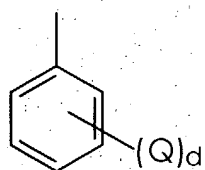
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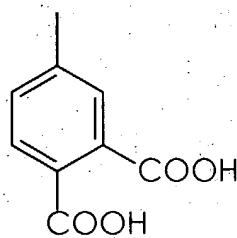
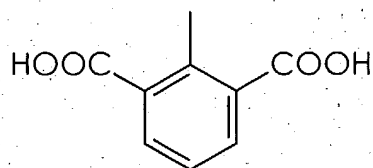
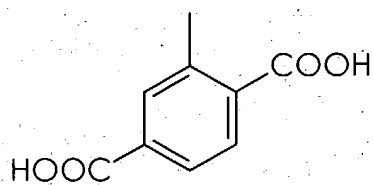
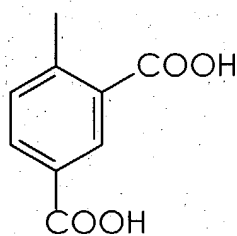
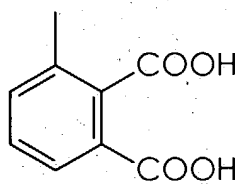
and



dicarboxylic acids and dicarboxylates, wherein

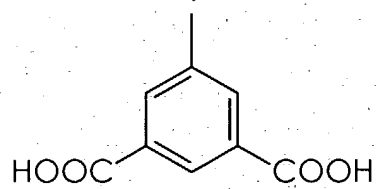


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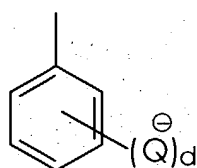


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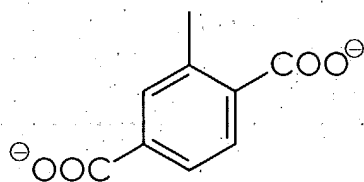
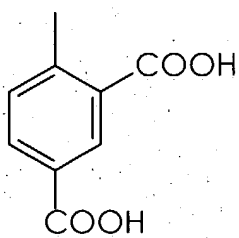
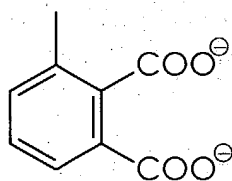
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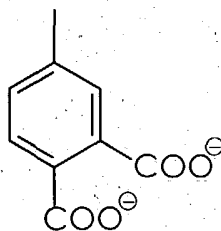
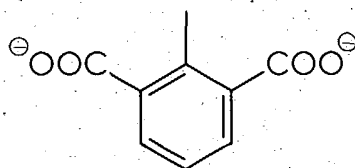


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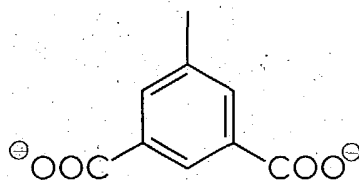


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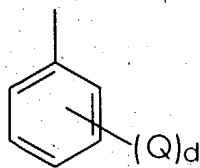




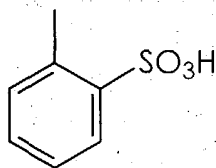
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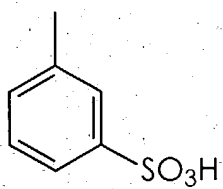


- 5 tricarboxylic acids and tricarboxylates, tetracarboxylic acids and tetracarboxylates, pentacarboxylic acids and pentacarboxylates, monosulfonic acids and monosulfonates, wherein

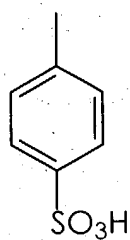


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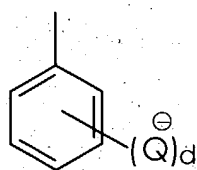




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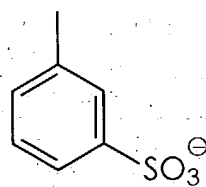
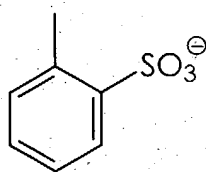


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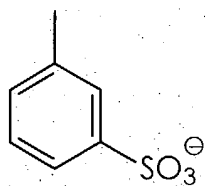


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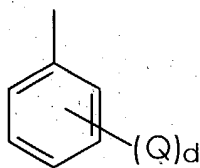
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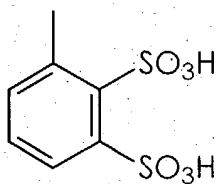
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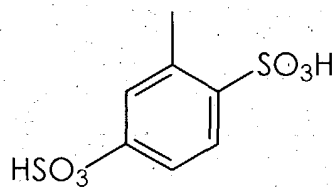
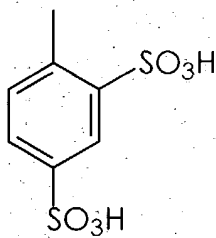
disulfonic acids and disulfonates, wherein

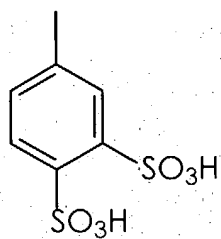
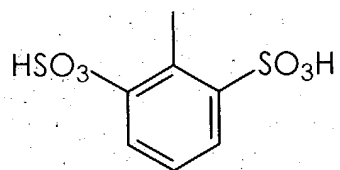


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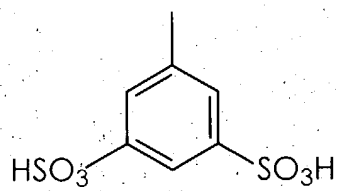


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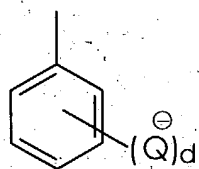




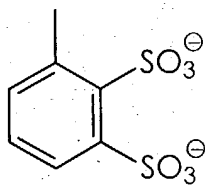
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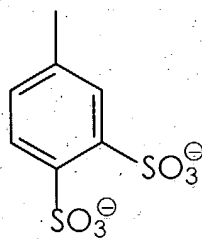
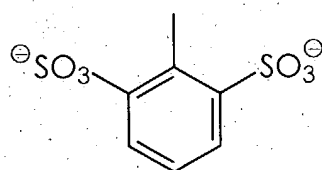
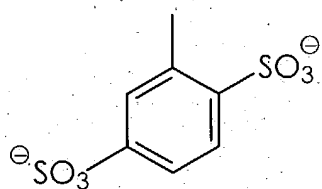
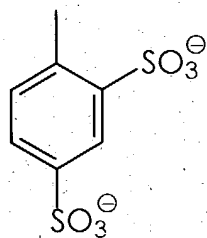


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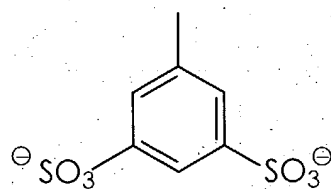


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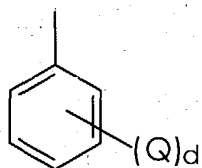




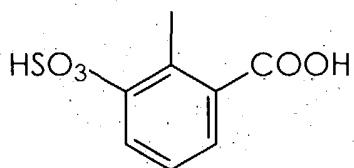
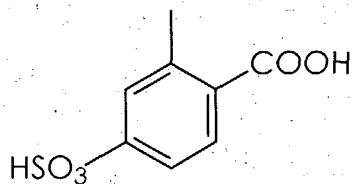
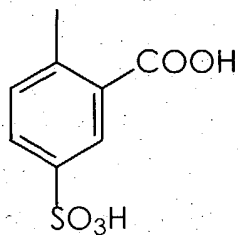
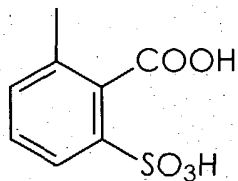
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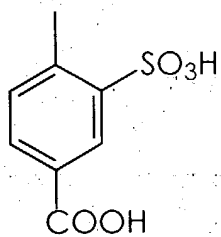
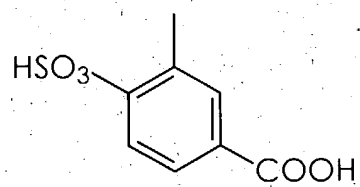
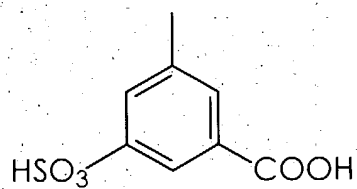
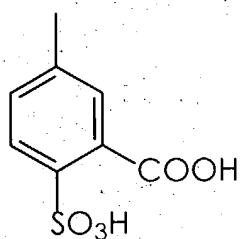
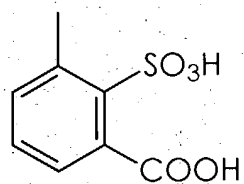


trisulfonic acids and trisulfonates, tetrasulfonic acids and tetrasulfonates, pentasulfonic acids and pentasulfonates, monocarboxylic acid monosulfonic acids and monocarboxylate monosulfonates, wherein



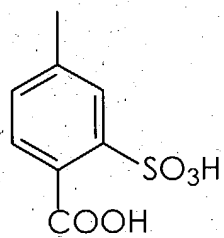
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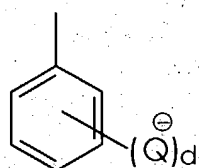


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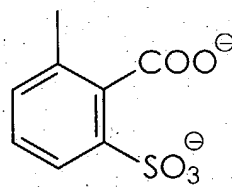
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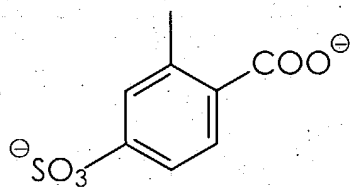
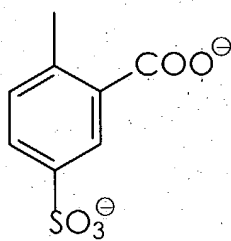
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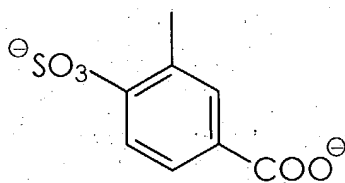
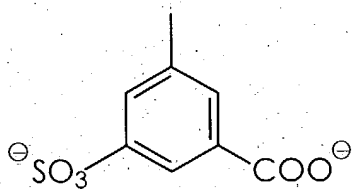
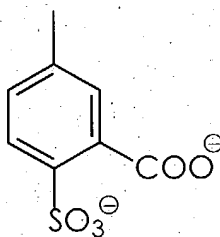
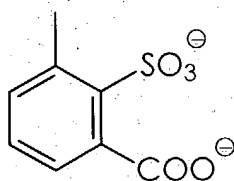
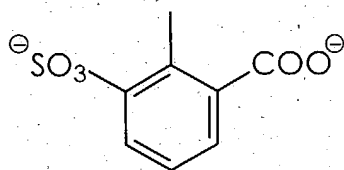


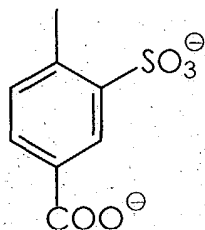
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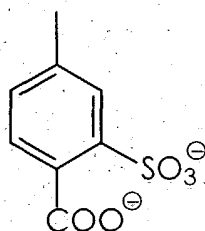
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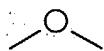


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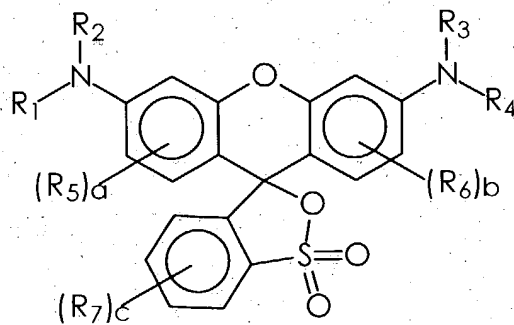
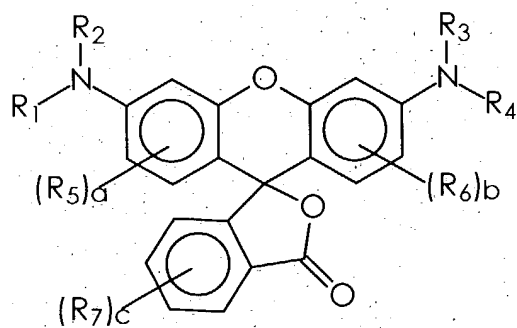


monocarboxylic acid disulfonic acids and monocarboxylate
 5 disulfonates, monocarboxylic acid trisulfonic acids and
 monocarboxylate trisulfonates, monocarboxylic acid tetrasulfonic acids
 and monocarboxylate tetrasulfonates, dicarboxylic acid monosulfonic
 acids and dicarboxylate monosulfonates, dicarboxylic acid disulfonic
 acids and dicarboxylate disulfonates, dicarboxylic acid trisulfonic acids
 10 and dicarboxylate trisulfonates, tricarboxylic acid monosulfonic acids
 and tricarboxylate monosulfonates, tricarboxylic acid disulfonic acids
 and tricarboxylate disulfonates, tetracarboxylic acid monosulfonic
 acids and tetracarboxylate monosulfonates, and the like. In addition, it
 is possible for a compound according to the present invention to have
 15 both one or more acid groups (i.e., COOH or SO₃H) and one or more
 anionic salt groups (i.e., COO⁻ or SO₃⁻) present in the molecule.

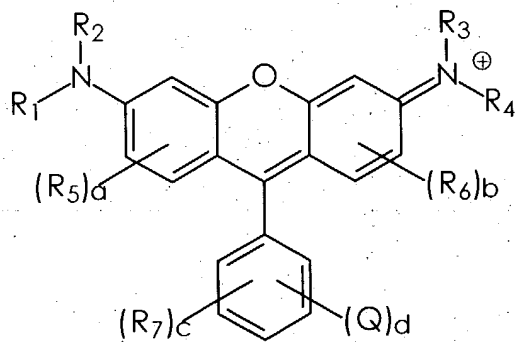
Colorant compounds suitable for inks according to the
 present invention include rhodamines, wherein



of the general formulae

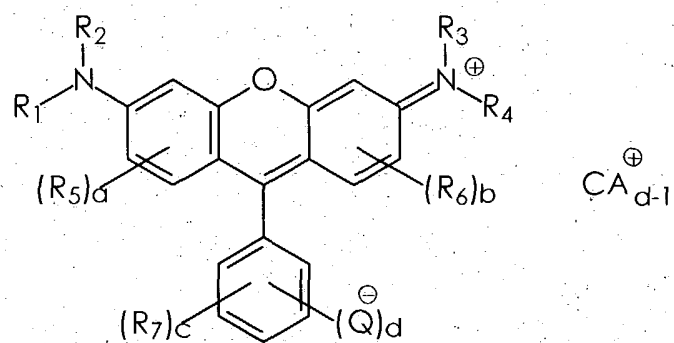


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A^{\ominus}

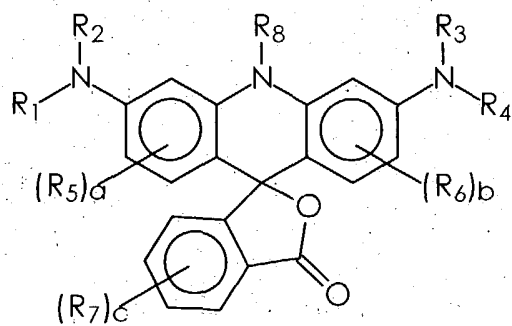
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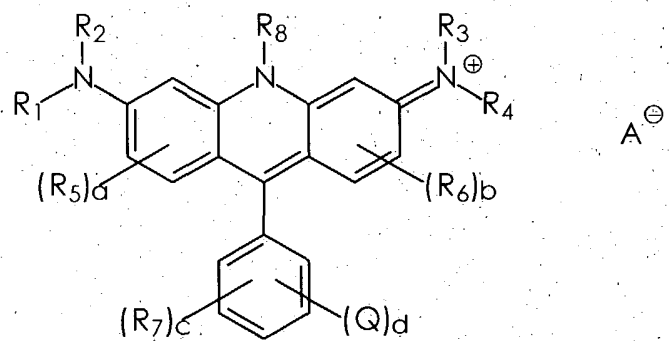
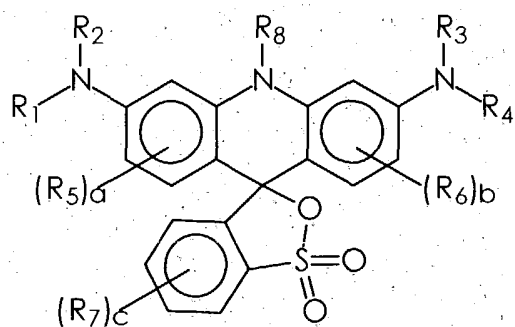


acridines, wherein

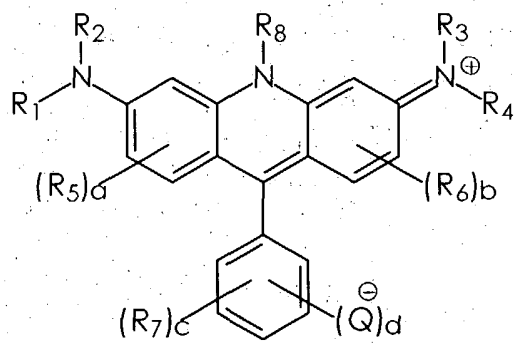


5 of the general formulae





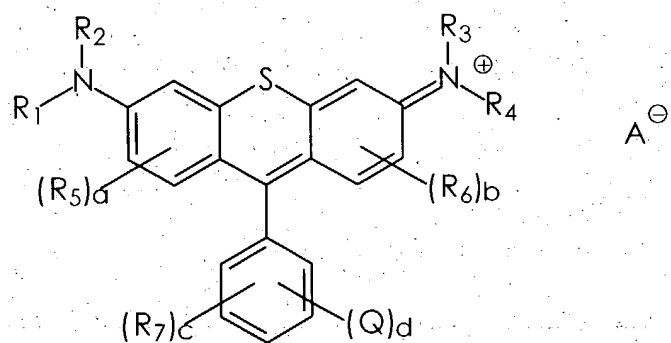
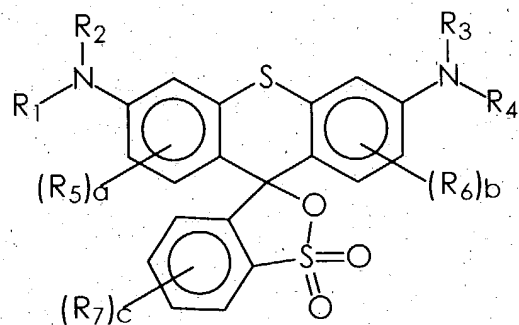
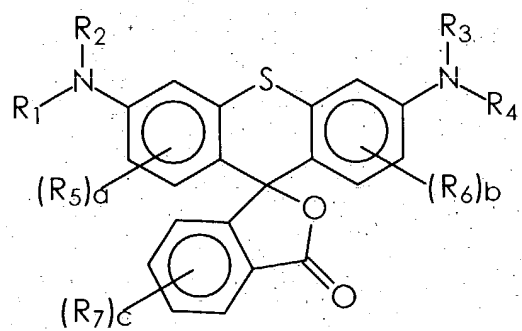
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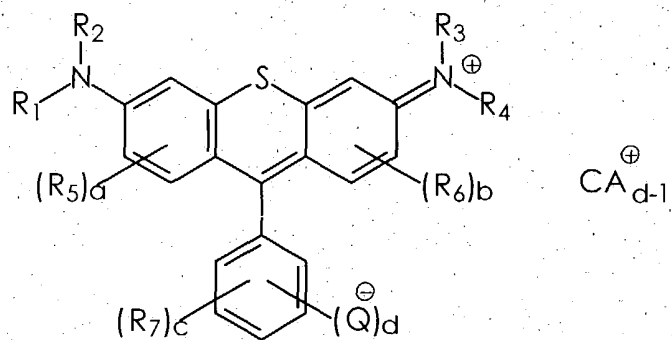
5 sulforhodamines, wherein

—Y— is

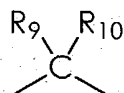
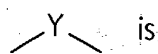
of the general formulae



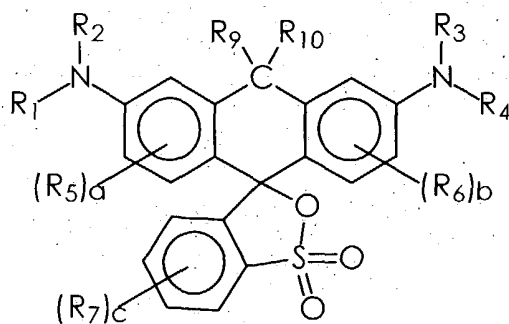
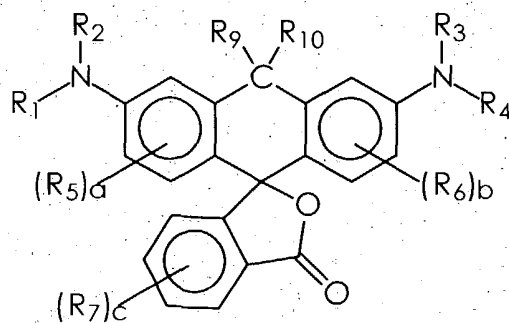
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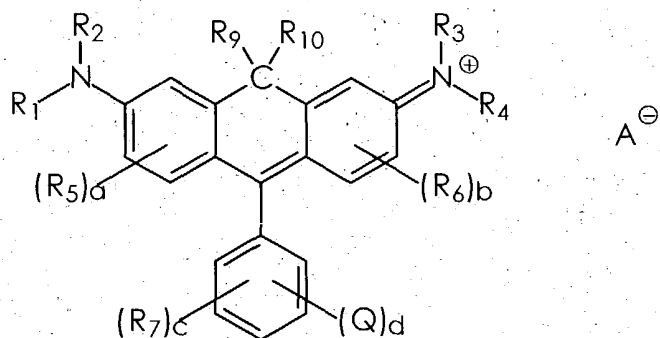


anthracenes, wherein

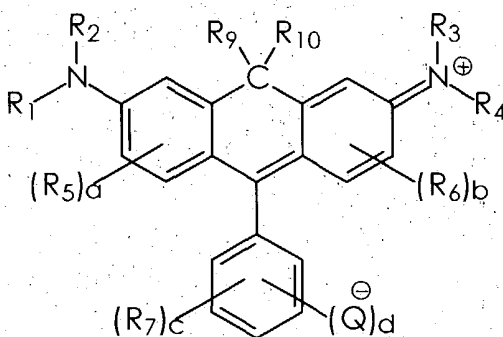


5 of the general formulae





and



and the like.

- 5 In a specific embodiment, the anion A can be an organic dianion of the formula $A_1-R_{11}-A_2$ wherein A_1 and A_2 each, independently of the other, are anionic groups, such as carboxylate, sulfonate, or the like, and wherein R_{11} is (i) an alkylene group (including linear, branched, saturated, unsaturated, cyclic, substituted, and
- 10 unsubstituted alkylene groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the alkylene group), in one embodiment with at least 1 carbon atom, in another embodiment with at least about 2 carbon atoms, in yet another embodiment with at least about 6 carbon

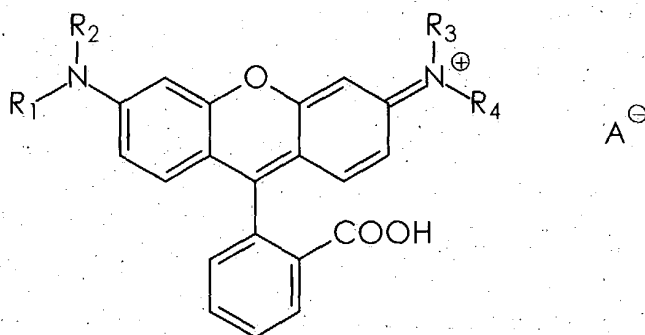
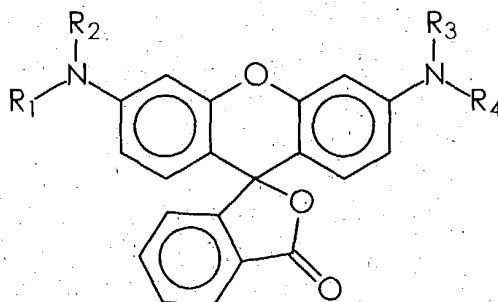
atoms, in another embodiment with at least about 8 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and
5 in yet another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, (ii) an arylene group (including unsubstituted and substituted arylene groups, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in the
10 arylene group), in one embodiment with at least about 6 carbon atoms, in another embodiment with at least about 10 carbon atoms, and in yet another embodiment with at least about 14 carbon atoms, and in one embodiment with no more than about 26 carbon atoms, in another embodiment with no more than about 22 carbon atoms, and
15 in yet another embodiment with no more than about 18 carbon atoms, although the number of carbon atoms can be outside of these ranges, (iii) an arylalkylene group (including unsubstituted and substituted arylalkylene groups, wherein the alkyl portion of the arylalkylene group can be linear, branched, saturated, unsaturated, and/or cyclic, and
20 wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the arylalkylene group), in one embodiment with at least about 7 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another
25 embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet

another embodiment with no more than about 20 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as benzyl or the like, or (iv) an alkylarylene group (including unsubstituted and substituted alkylarylene groups, wherein the alkyl
5 portion of the alkylarylene group can be linear, branched, saturated, unsaturated, and/or cyclic, and wherein hetero atoms, such as oxygen, nitrogen, sulfur, silicon, phosphorus, and the like either may or may not be present in either or both of the alkyl portion and the aryl portion of the alkylarylene group), in one embodiment with at least about 7
10 carbon atoms, in another embodiment with at least about 12 carbon atoms, and in yet another embodiment with at least about 18 carbon atoms, and in one embodiment with no more than about 55 carbon atoms, in another embodiment with no more than about 30 carbon atoms, and in yet another embodiment with no more than about 20
15 carbon atoms, although the number of carbon atoms can be outside of these ranges, such as tolyl or the like, and wherein the substituents on the substituted alkylene, arylene, arylalkylene, and alkylarylene groups can be (but are not limited to) hydroxy groups, halogen atoms, amine groups, imine groups, ammonium groups, cyano groups, pyridine
20 groups, pyridinium groups, ether groups, aldehyde groups, ketone groups, ester groups, amide groups, carbonyl groups, thiocarbonyl groups, sulfate groups, sulfonate groups, sulfonic acid groups, sulfide groups, sulfoxide groups, phosphine groups, phosphonium groups, phosphate groups, nitrile groups, mercapto groups, nitro groups, nitroso
25 groups, sulfone groups, acyl groups, acid anhydride groups, azide groups, azo groups, cyanato groups, isocyanato groups, thiocyanato groups, isothiocyanato groups, carboxylate groups, carboxylic acid

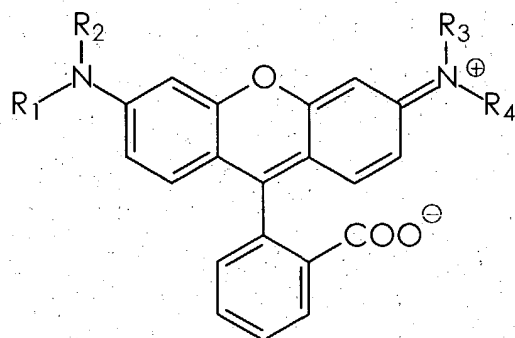
groups, urethane groups, urea groups, mixtures thereof, and the like, wherein two or more substituents can be joined together to form a ring. Examples of suitable organic dianions include unsubstituted and substituted naphthalene disulfonates, unsubstituted and substituted
 5 benzene disulfonates, and the like, as well as mixtures thereof.

In another specific embodiment, the anion A can be an organic trianion, tetraanion, and higher, an oligomeric and polymeric anion, such as a polysulfonate or polycarboxylate, or the like.

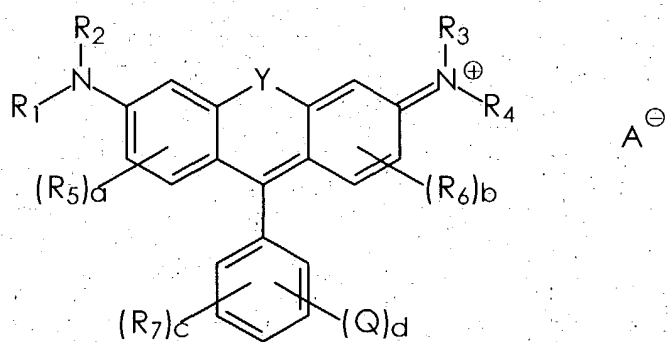
In one specific embodiment, the colorant compounds for
 10 inks according to the present invention are of the formulae



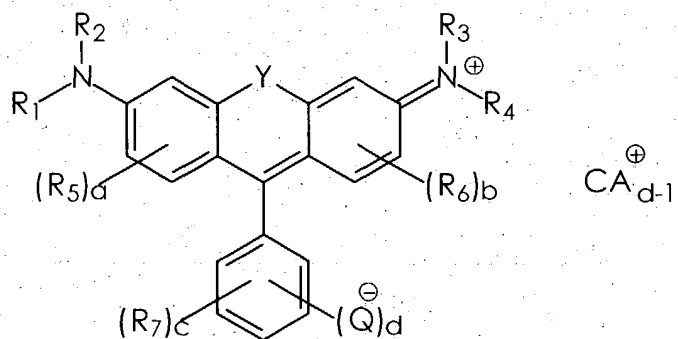
and



It is to be understood that in colorants of the formulae

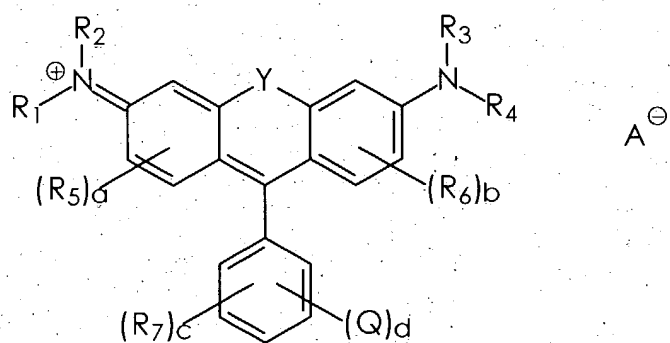
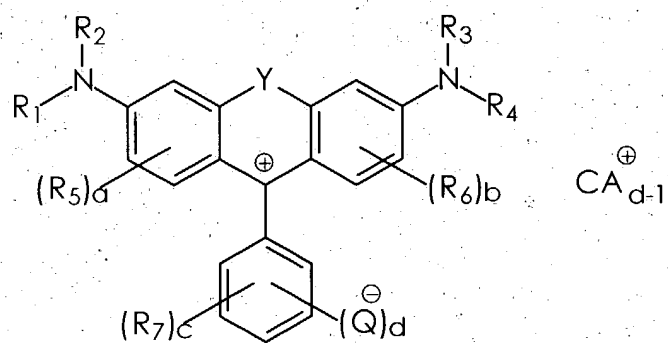
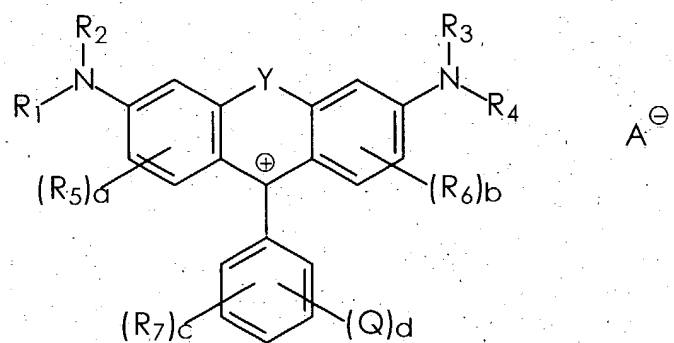


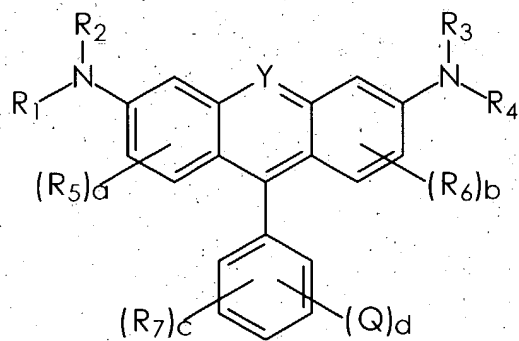
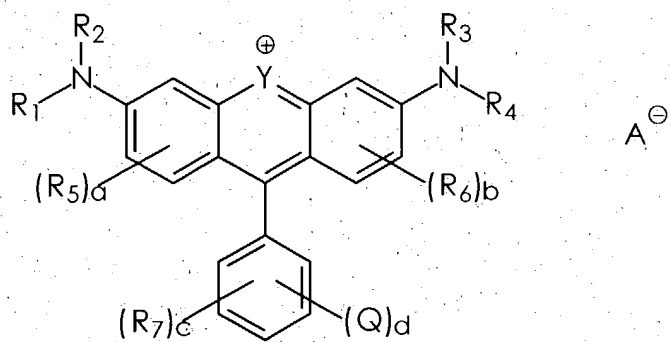
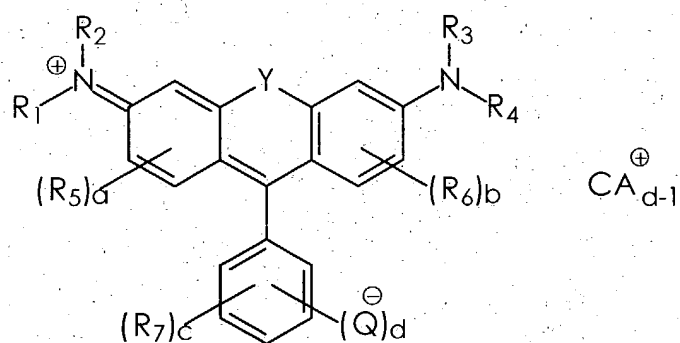
and



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the positive charge is delocalized, and that other tautomeric structures can be drawn, including (but not limited to)





and the like. It is to be understood that all possible tautomeric forms of
 5 these colorants are included within the above formulae.

Colorant compounds for inks of the present invention can be prepared by any desired or effective procedure. For example, a

dihalofluorescein, such as dichlorofluorescein or the like, can be admixed with one or more amines having the desired R₁, R₂, R₃, and R₄ groups thereon, an optional zinc halide, such as zinc chloride or the like, and an optional nonnucleophilic base, such as calcium oxide, zinc
5 oxide, or the like, either neat or, optionally, in the presence of a solvent.

The amine and the dihalofluorescein are present in any desired or effective relative amounts, in one embodiment at least about 0.9 mole of base per every one mole of dihalofluorescein, in another embodiment at least about 0.95 mole of base per every one
10 mole of dihalofluorescein, and in yet another embodiment at least about 1 mole of base per every one mole of dihalofluorescein, and in one embodiment no more than about 20 moles of base per every one mole of dihalofluorescein, in another embodiment no more than about 10 moles of base per every one mole of dihalofluorescein, and in yet
15 another embodiment no more than about 2 moles of base per every one mole of dihalofluorescein, although the relative amounts can be outside of these ranges.

Dichlorofluorescein is commercially available from, for example, Aldrich Chemical Co., Milwaukee, WI. Dihalofluoresceins can
20 also be prepared by the reaction of fluorescein with PX₅ wherein X is fluorine, chlorine, bromine, or iodine, or with a toluenesulfonylhalide, such as toluenesulfonylchloride or the like.

When an optional zinc halide is used, the dihalofluorescein and the zinc halide are present in any desired or effective relative
25 amounts, in one embodiment at least about 2 moles of zinc halide per every one mole of dihalofluorescein, in another embodiment at least about 2.5 moles of zinc halide per every one mole of dihalofluorescein,

and yet in another embodiment at least about 3 moles of zinc halide per every one mole of dihalofluorescein, and in one embodiment no more than about 5 moles of zinc halide per every one mole of dihalofluorescein, in another embodiment no more than about 4.5 moles of zinc halide per every one mole of dihalofluorescein, and in yet another embodiment no more than about 4 moles of zinc halide per every one mole of dihalofluorescein, although the relative amounts can be outside of these ranges.

When an optional base is used, the base is present in any desired or effective amount, in one embodiment at least about 2 equivalents of base per every one mole of dihalofluorescein (i.e., about 2 moles of monobasic base per every one mole of dihalofluorescein, about 1 mole of dibasic base, such as calcium oxide, per every one mole of dihalofluorescein, and the like), in another embodiment at least about 2.5 equivalents of base per every one mole of dihalofluorescein, and yet in another embodiment at least about 3 equivalents of base per every one mole of dihalofluorescein, and in one embodiment no more than about 10 equivalents of base per every one mole of dihalofluorescein, in another embodiment no more than about 5 equivalents of base per every one mole of dihalofluorescein, and in yet another embodiment no more than about 3.2 equivalents of base per every one mole of dihalofluorescein, although the relative amounts can be outside of these ranges.

If desired, the reaction can be run neat, in the absence of a solvent. In addition, if desired, the reaction can be run in the presence of an optional solvent. Examples of suitable solvents include tetramethylene sulfone (sulfolane), N-methyl pyrrolidone, dimethyl

formamide, dimethyl sulfoxide, octanol, or the like, as well as mixtures thereof. When present, the optional solvent is present in any desired or effective amount, in one embodiment at least about 1 liter per every 0.1 mole of dihalofluorescein, in another embodiment at least about 1 liter per every 0.3 mole of dihalofluorescein, and in yet another embodiment at least about 1 liter per every 0.35 mole of dihalofluorescein, and in one embodiment no more than about 1 liter per every 2 moles of dihalofluorescein, in another embodiment no more than about 1 liter per every 1.5 moles of dihalofluorescein, and in yet another embodiment no more than about 1 liter per every 1 mole of dihalofluorescein, although the relative amounts can be outside of these ranges.

The mixture of dihalofluorescein, amine, optional zinc halide, optional base, and optional solvent is then heated to any effective temperature, in one embodiment at least about 62°C, in another embodiment at least about 150°C, and in yet another embodiment at least about 190°C, and in one embodiment no more than about 280°C, in another embodiment no more than about 220°C, and in yet another embodiment no more than about 200°C, although the temperature can be outside of these ranges.

The mixture of dihalofluorescein, amine, optional zinc halide, optional base, and optional solvent is heated for any effective period of time, in one embodiment at least about 5 minutes, in another embodiment at least about 2 hours, and in yet another embodiment at least about 3 hours, and in one embodiment no more than about 4 days, in another embodiment no more than about 60 hours, and in yet

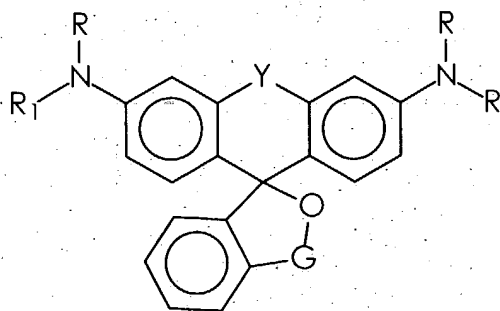
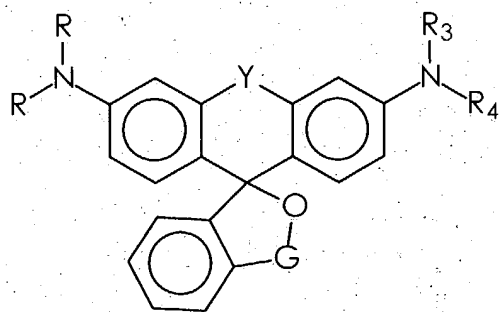
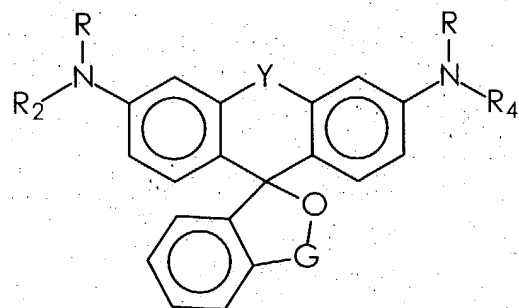
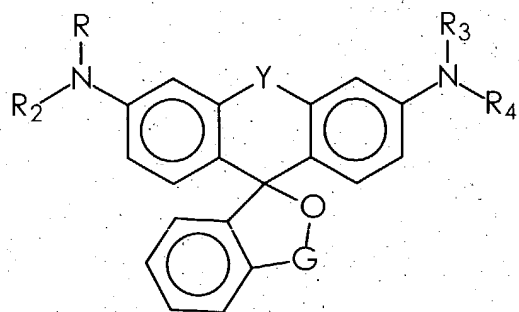
another embodiment no more than about 40 hours, although the time can be outside of these ranges.

If desired, the resulting product can be purified by pouring the reaction mixture into an organic non-water-soluble and non-water-miscible solvent in which the product is soluble or miscible and in which undesirable salt byproducts are not soluble, such as methyl isobutyl ketone, toluene, hexane, heptane, or the like, followed by admixing the solvent containing the product with water in a separatory funnel and separating the aqueous and organic phases.

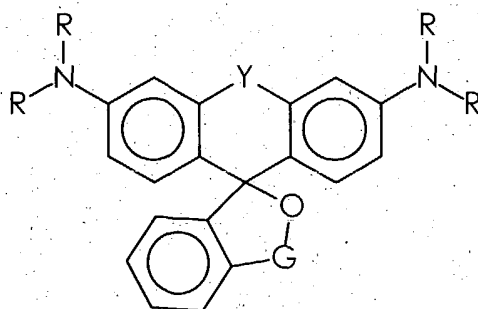
The crude product can then, if desired, be further purified by washing it with aqueous EDTA to remove metal salts, followed by washing with water. If desired, a titration or other instrumental technique, such as AA (atomic absorption) or ICP (inductively coupled plasma) can be performed to determine if the metal salts have been completely removed. The purified product can be isolated by distilling off any solvents.

Various substituents can be placed on the rings of the colorants of the present invention by any desired or effective method, such as, for example, the methods disclosed in U.S. Patent 5,847,162 and U.S. Patent 1,991,482, the disclosures of each of which are totally incorporated herein by reference.

Additional numbers of carbon atoms can be placed on the central structure by, for example, selecting long chain amines as reactants. Examples of such compounds include (but are not limited to) those of the formulae



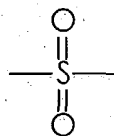
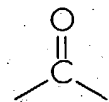
and



wherein Y, R₁, R₂, R₃, and R₄ have the same definitions as given hereinabove, G is either

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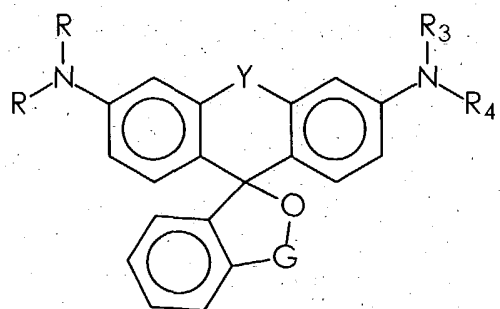
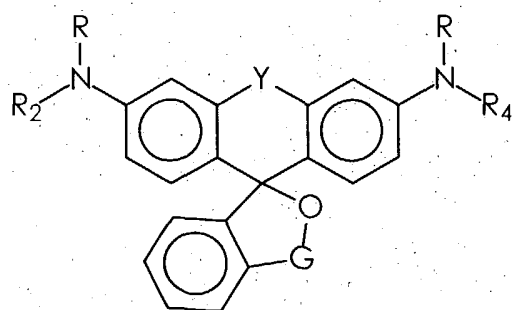
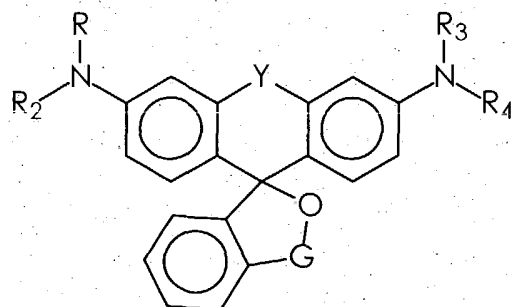
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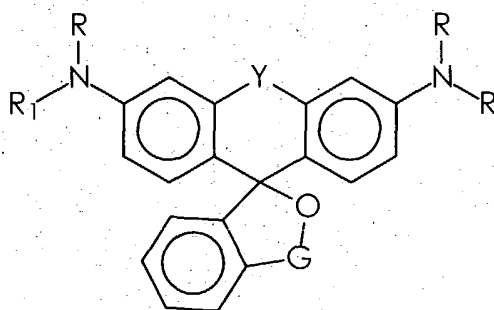


and (1) R is a linear alkyl group of the formula -C_nH_{2n+1} wherein n is at least about 12, (2) R is a branched alkyl group of the formula -C_nH_{2n+1} wherein n is at least about 12, (3) R is an ether group of the formula -(CH₂)₃-O-C_nH_{2n+1} wherein n is at least about 11, and the like, as well as their ring-opened, or protonated, or free-base forms and their zwitterionic forms.

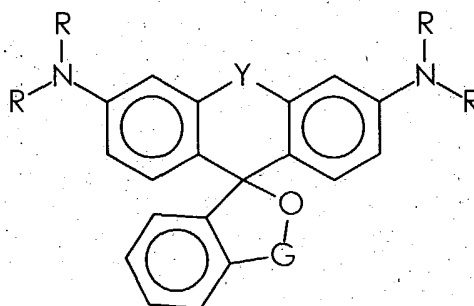
Additional numbers of carbon atoms can also be placed on the central structure by, for example, first preparing the corresponding alcohols and then reacting these alcohols with, for example, high-carbon-number acids to prepare esters, high-carbon-

number isocyanates to prepare urethanes, or the like. Examples of such compounds include (but are not limited to) those of the formulae

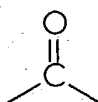




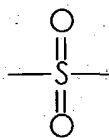
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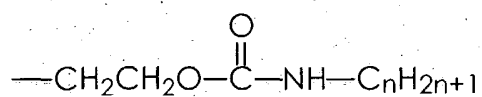
wherein Y, R₁, R₂, R₃, and R₄ have the same definitions as given
 5 hereinabove, G is either



or

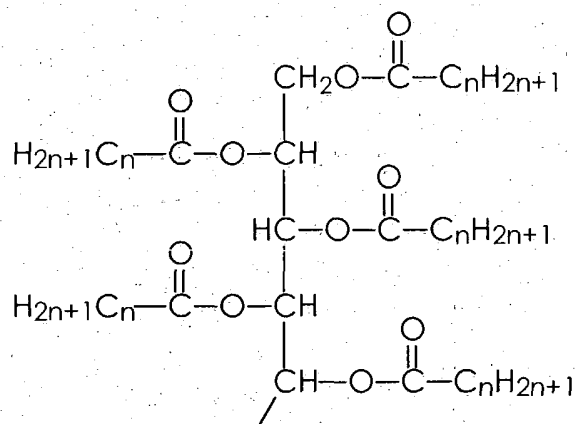


and (1) R is a group of the formula

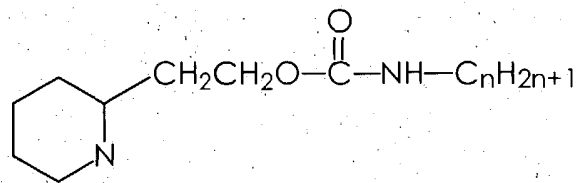


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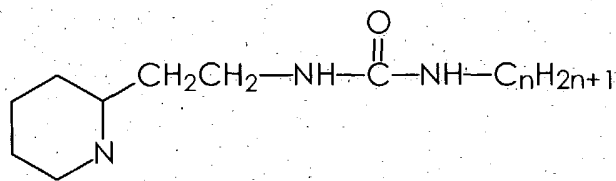
wherein n is at least about 12, (2) R is a group of the formula



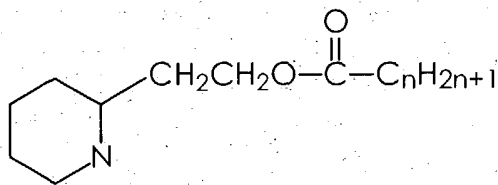
wherein n is at least about 12, (7) two R groups on the same nitrogen atom form a group, with the nitrogen atom, of the formula



- 5 wherein n is at least about 12, (8) two R groups on the same nitrogen atom form a group, with the nitrogen atom, of the formula

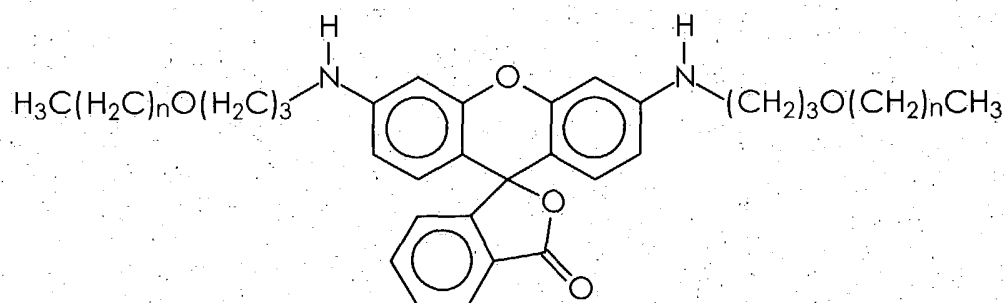


wherein n is at least about 12, (9) two R groups on the same nitrogen atom form a group, with the nitrogen atom, of the formula

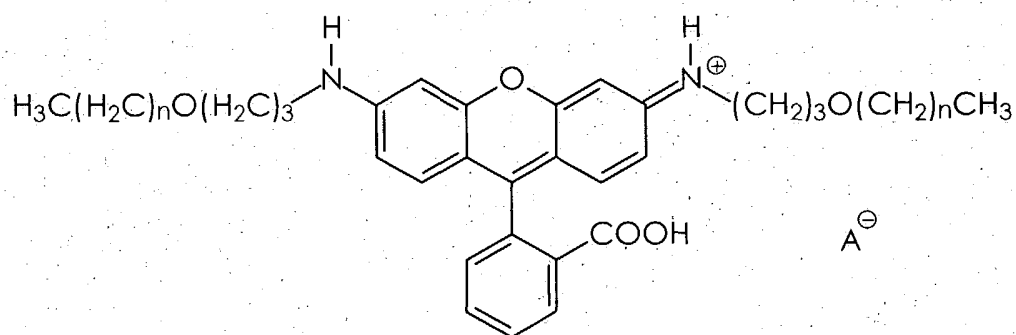


wherein n is at least about 12, and the like, as well as their ring-opened, or protonated, or free-base forms and their zwitterionic forms.

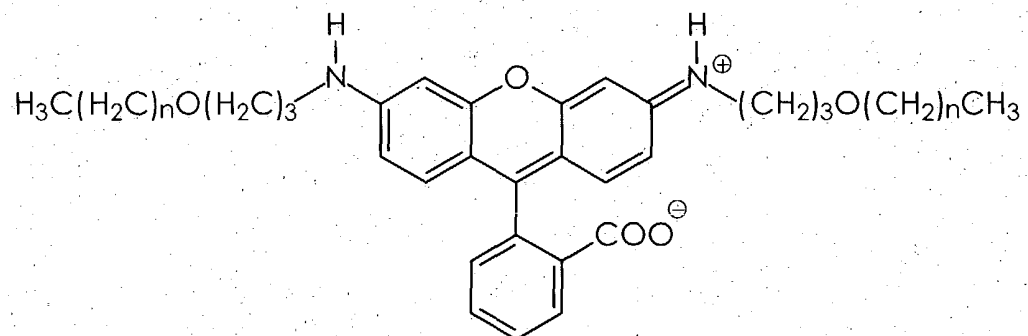
Some specific examples of such compounds include (a) those of the formulae



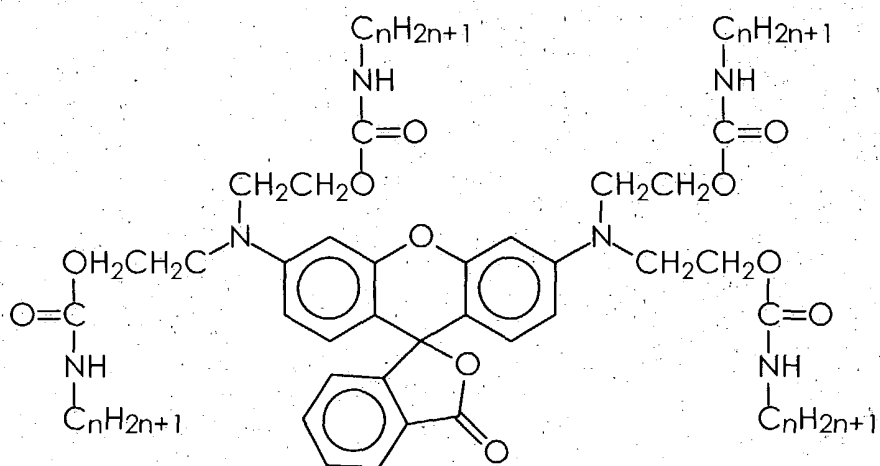
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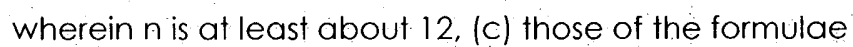


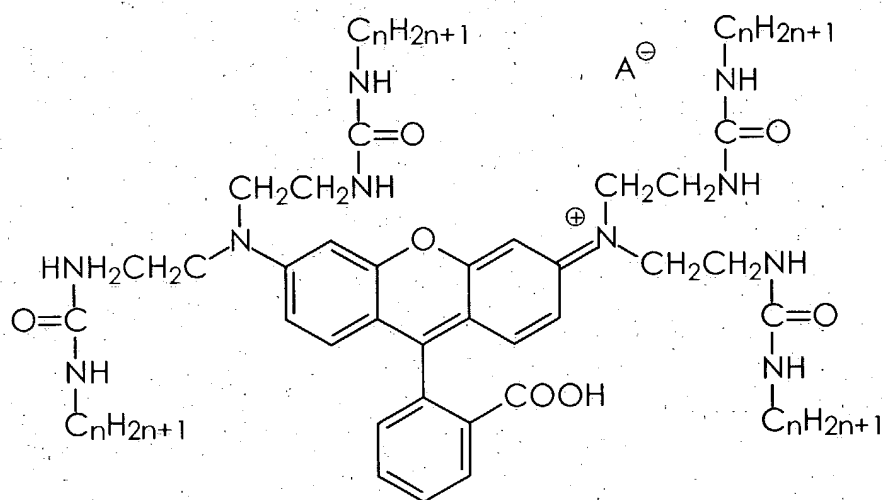
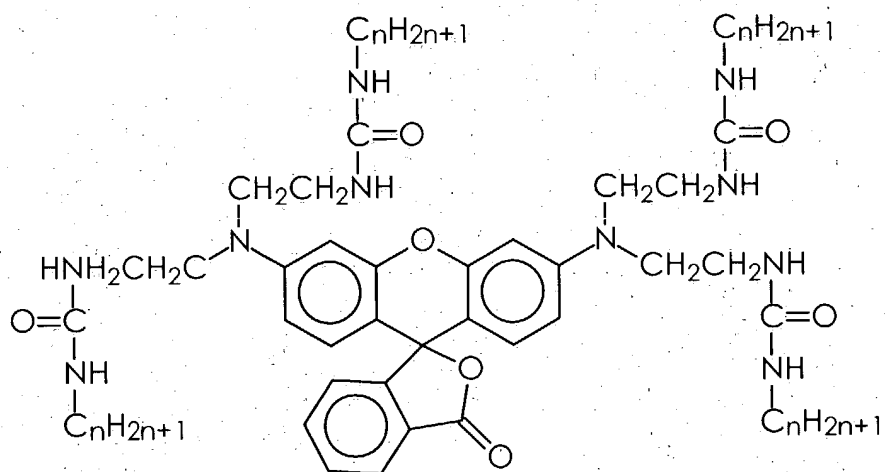
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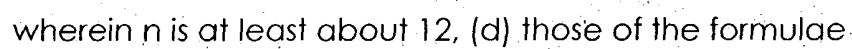
wherein n is at least about 11, (b) those of the formulae

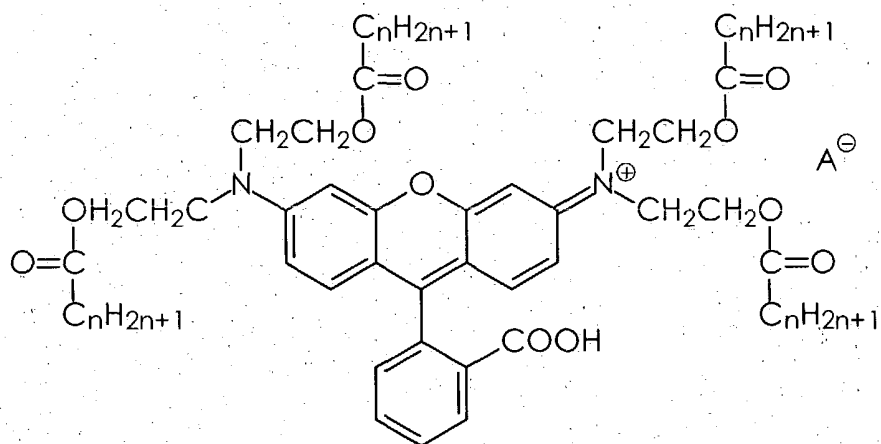




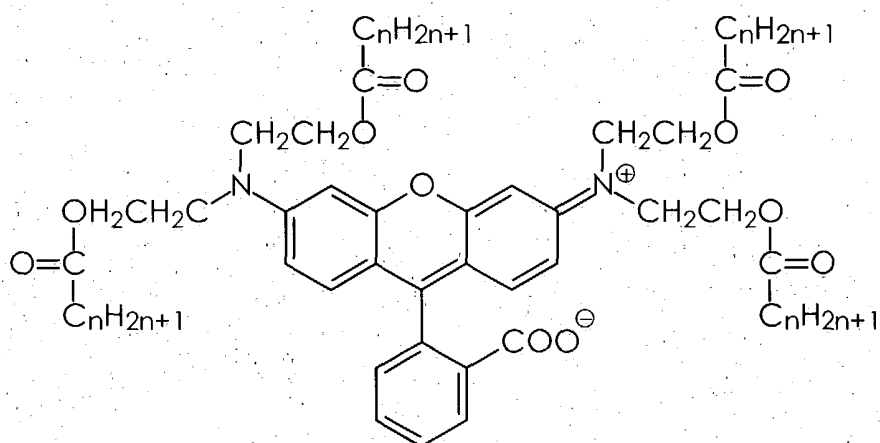


and

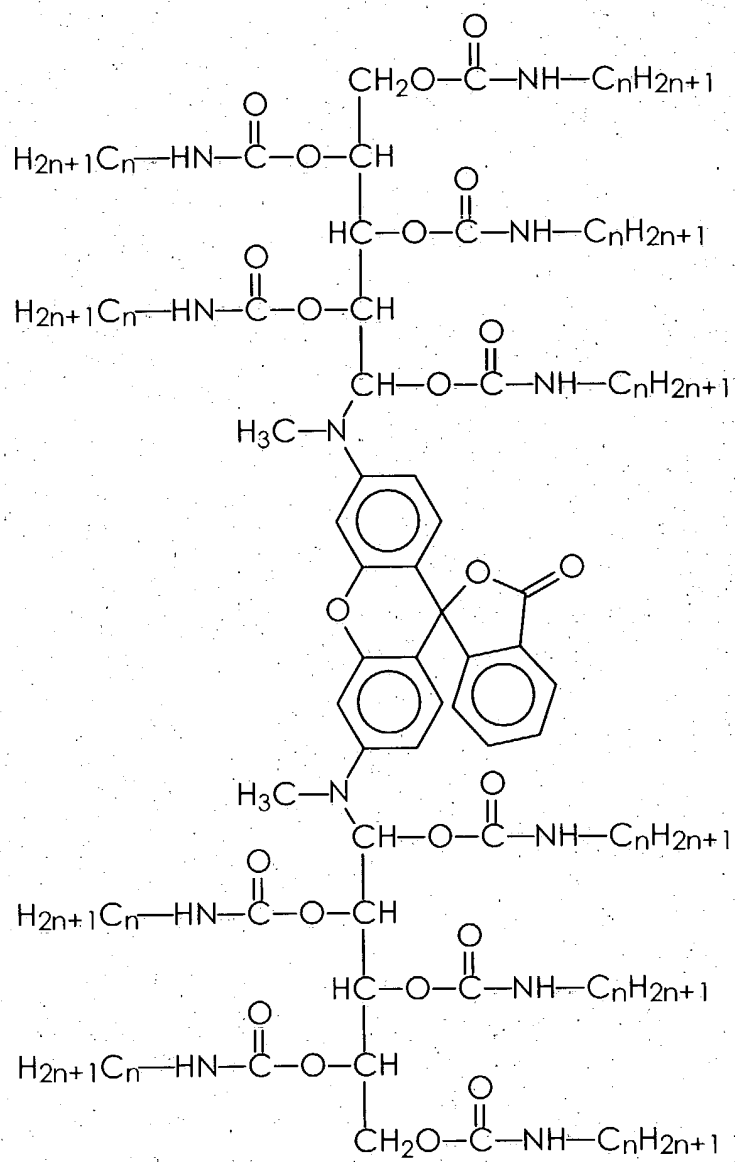


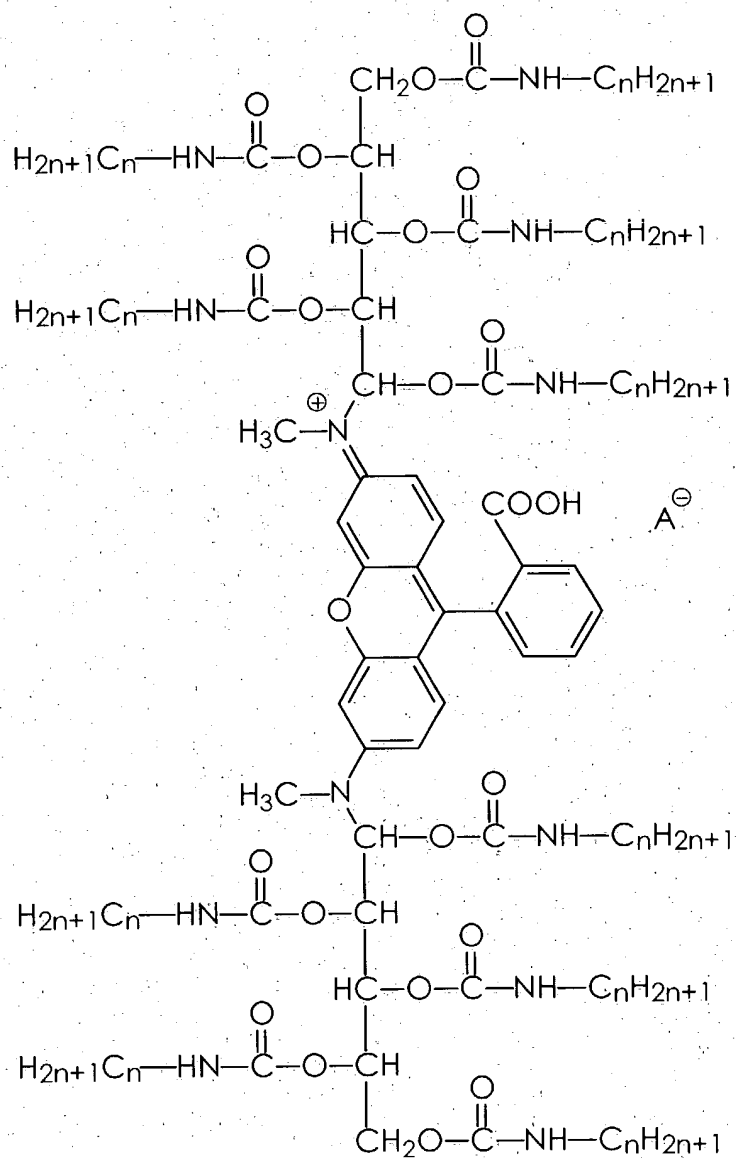


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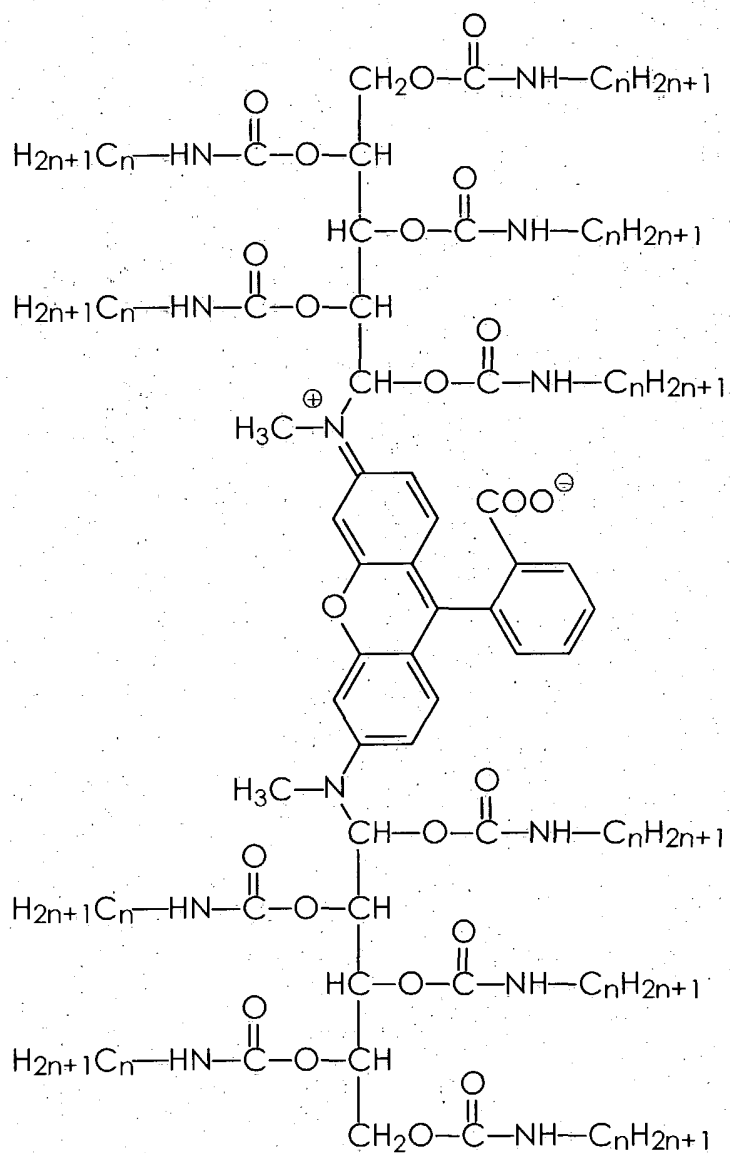


wherein n is at least about 12, (e) those of the formulae

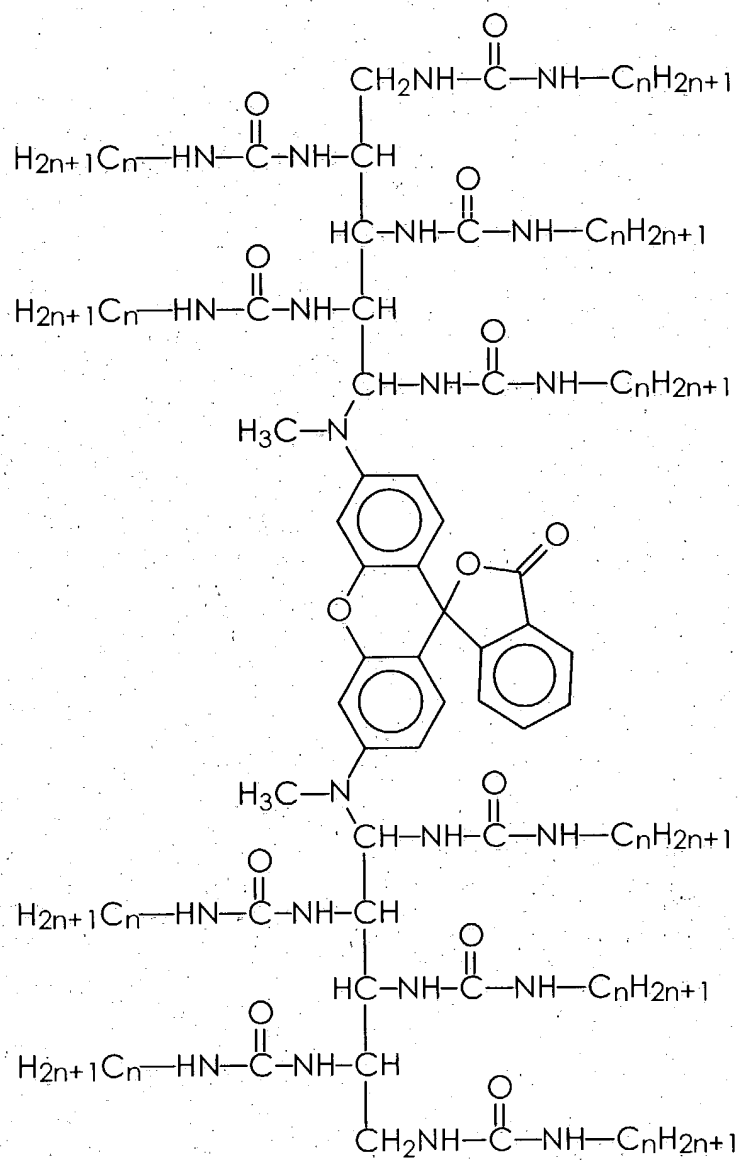


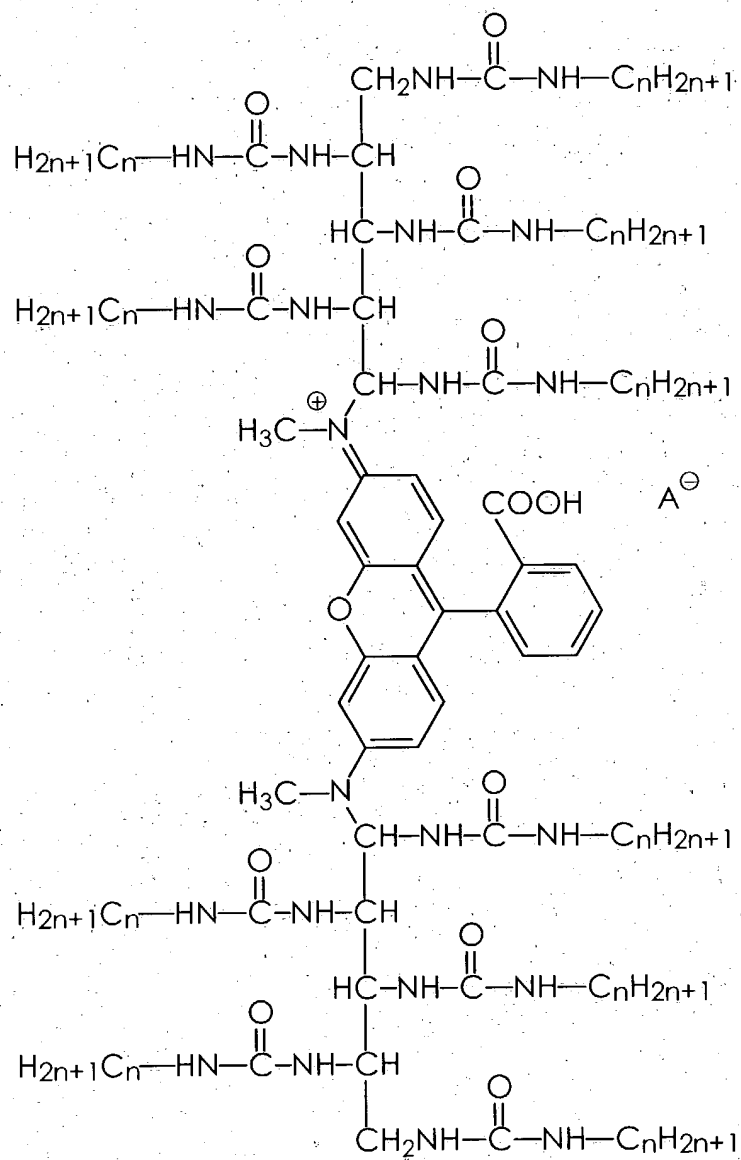


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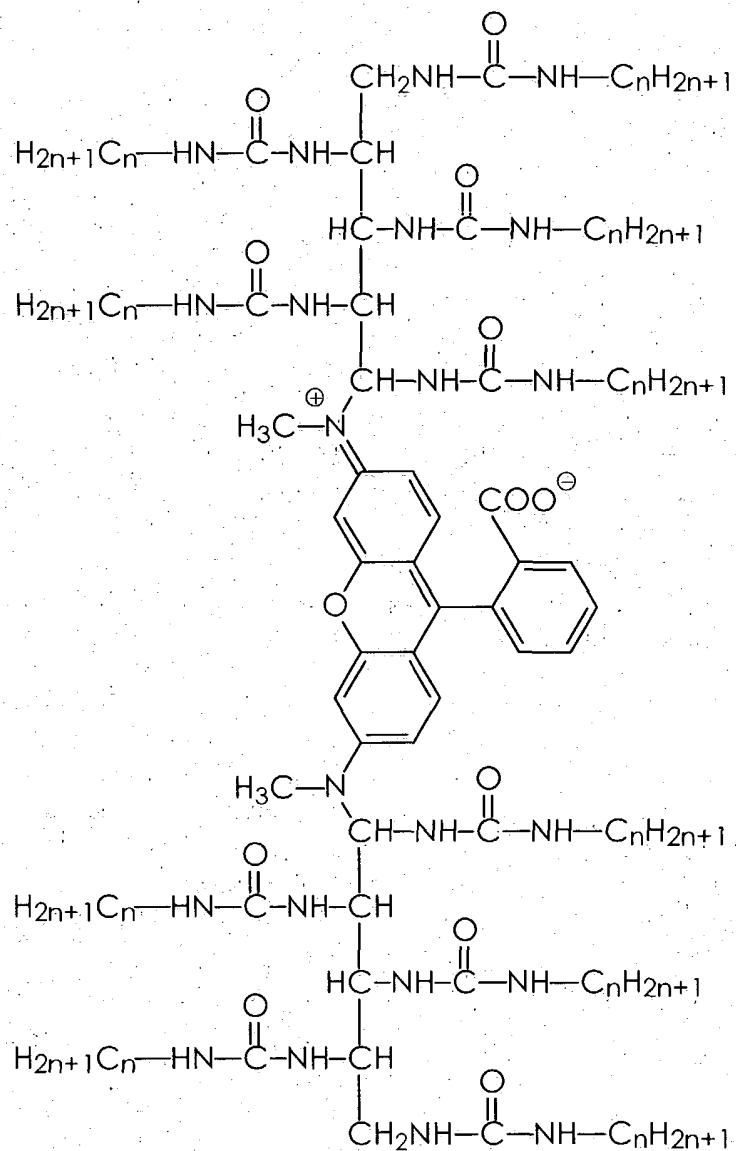


wherein n is at least about 12, (f) those of the formulae

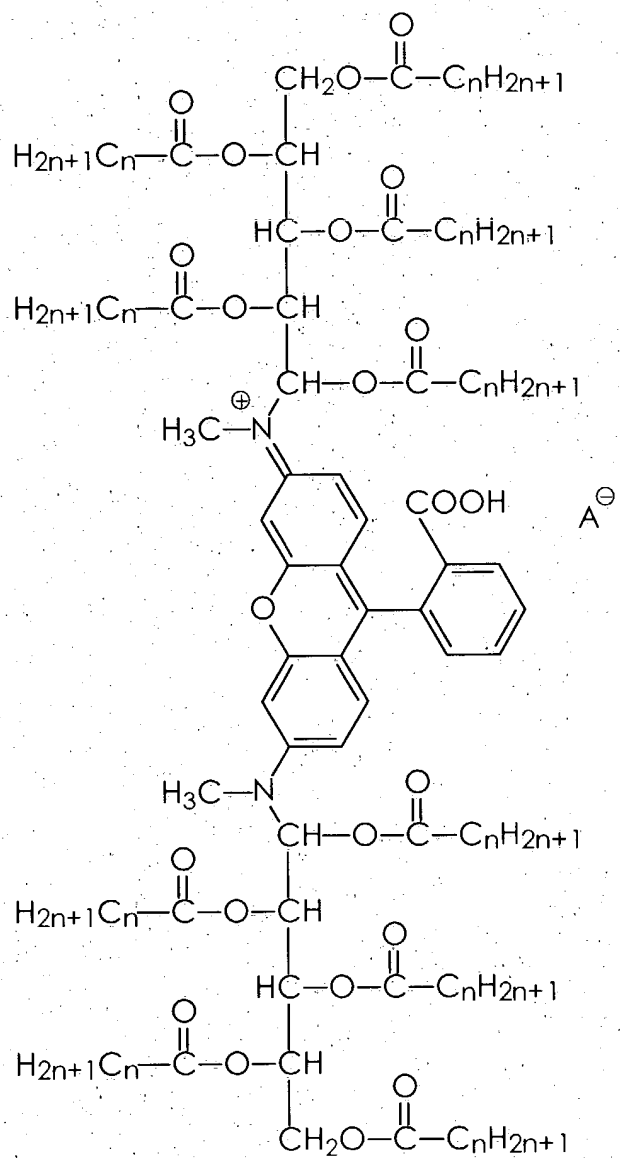




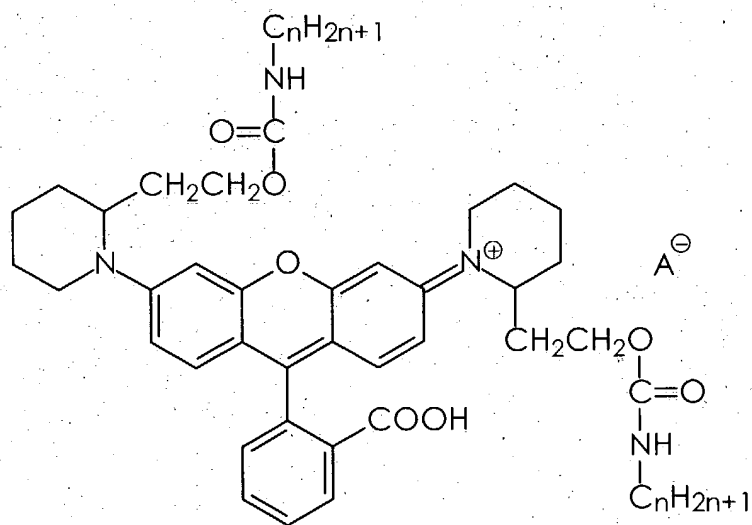
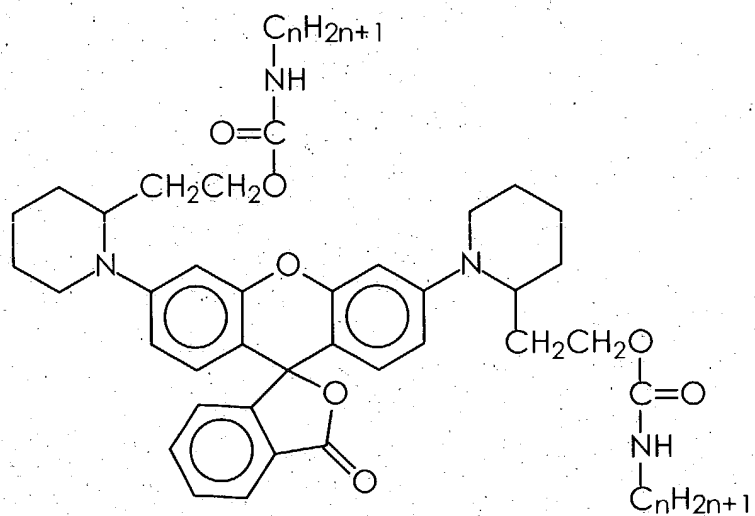
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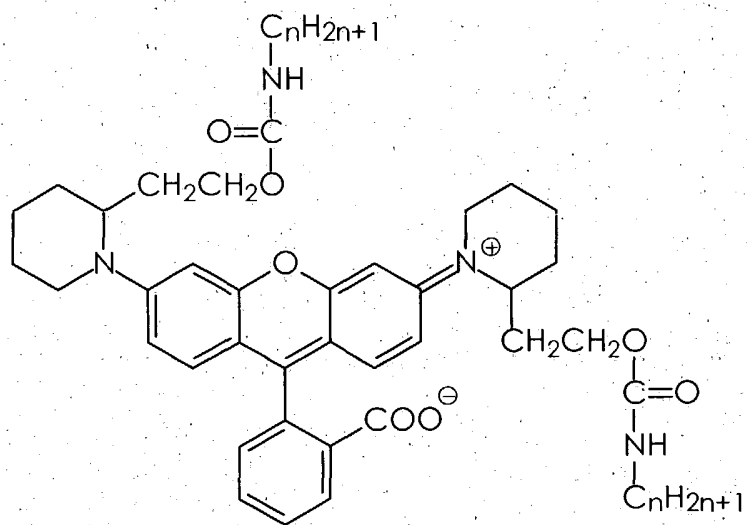
wherein n is at least about 12, (g) those of the formulae



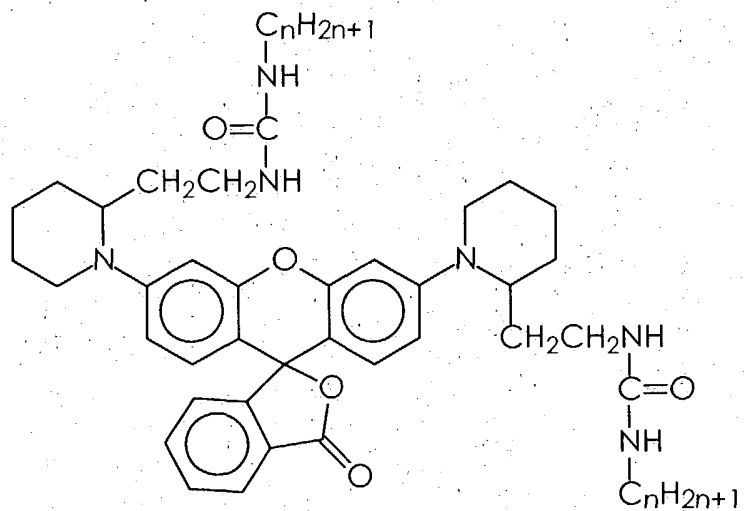
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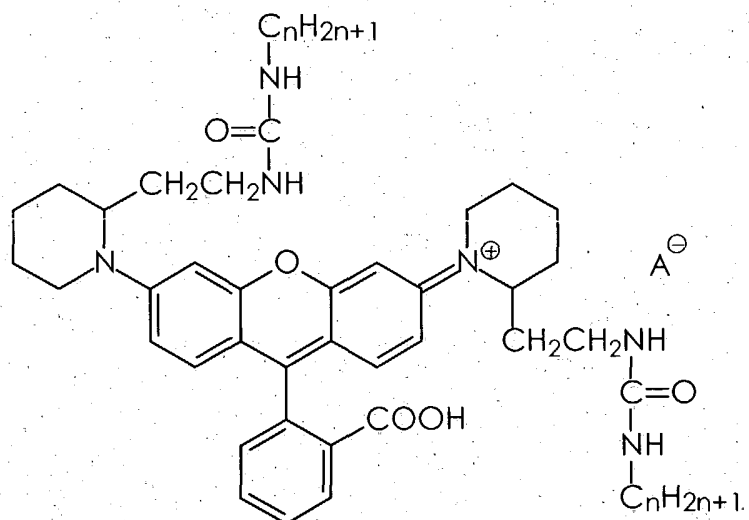


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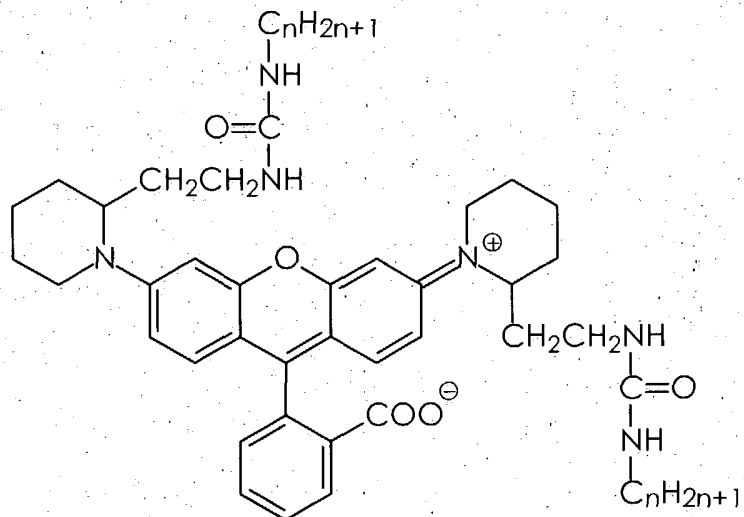


wherein n is at least about 12, (i) those of the formulae

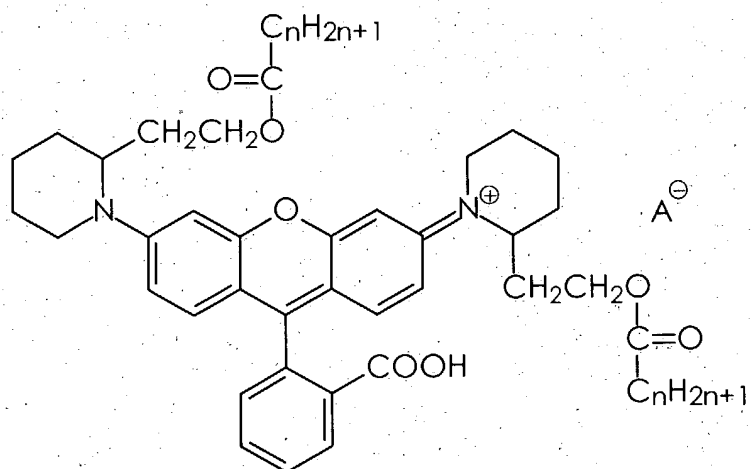
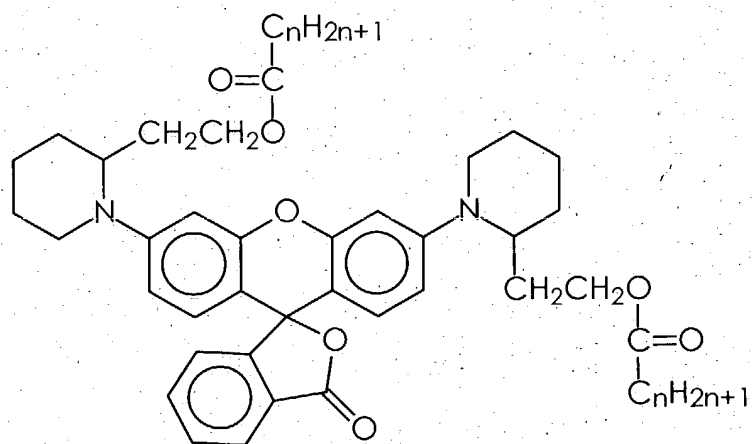




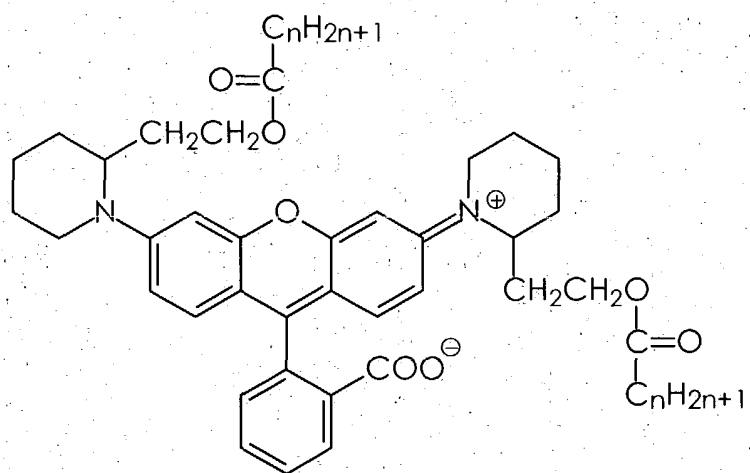
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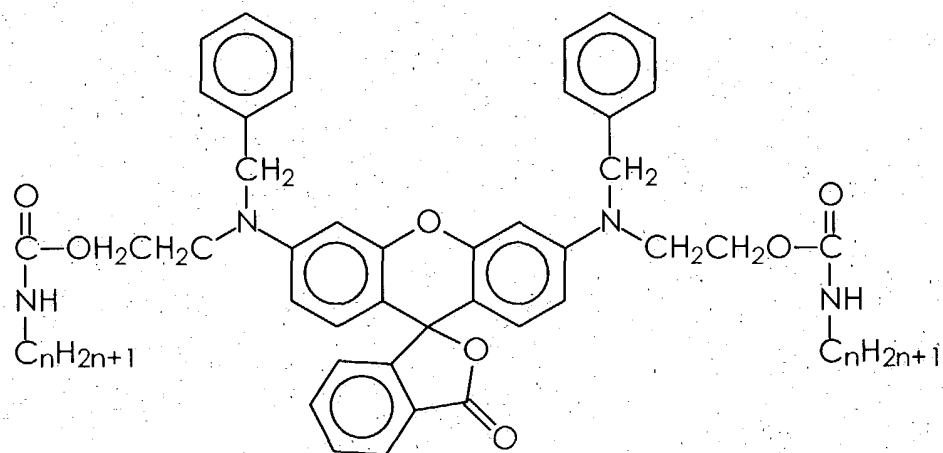
wherein n is at least about 12, (j) those of the formulae

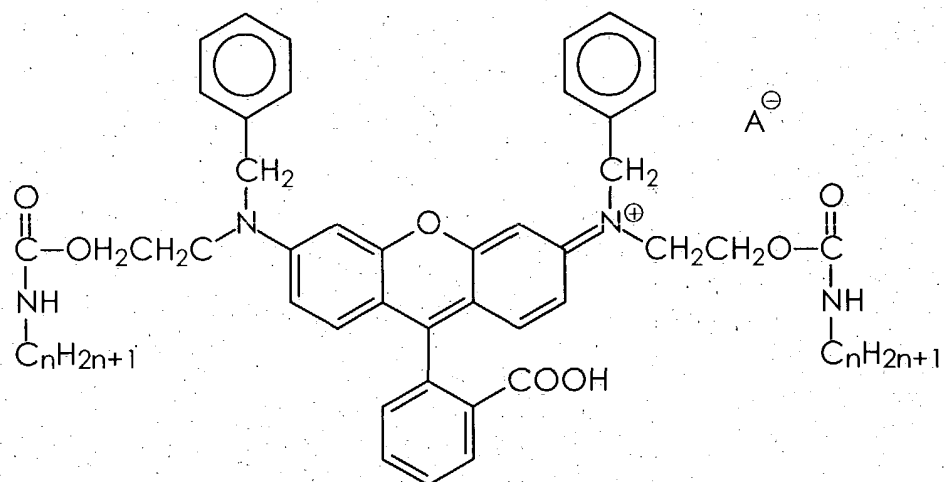


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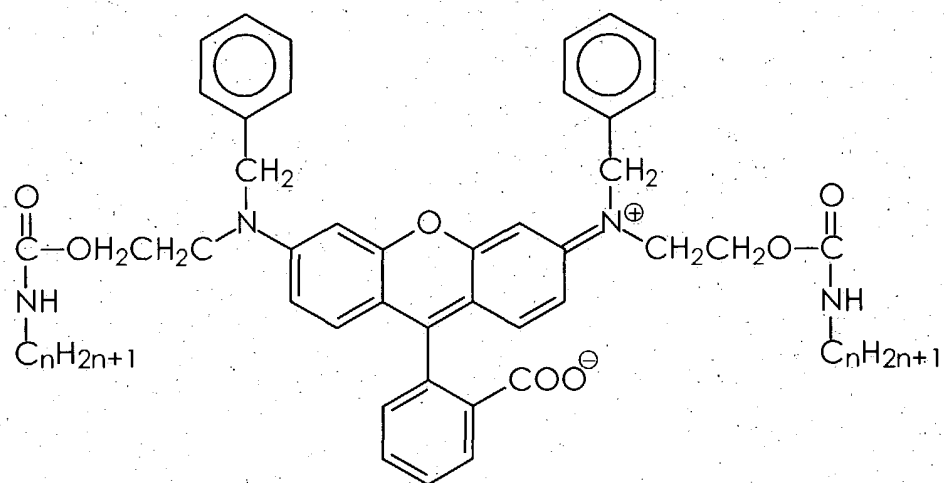


wherein n is at least about 12, (k) those of the formulae

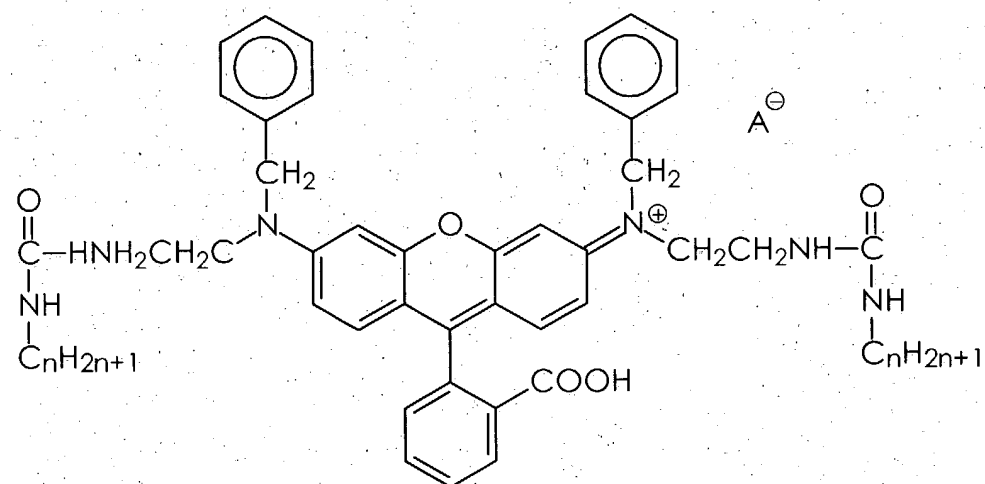
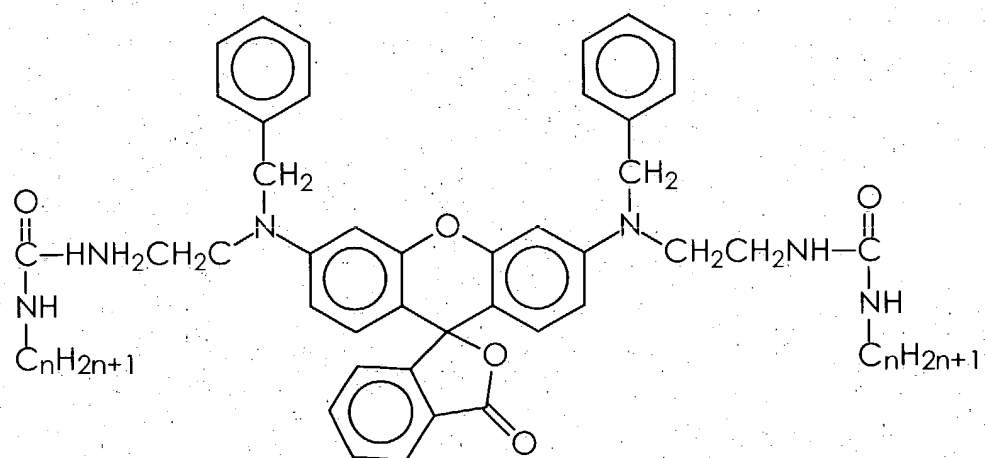




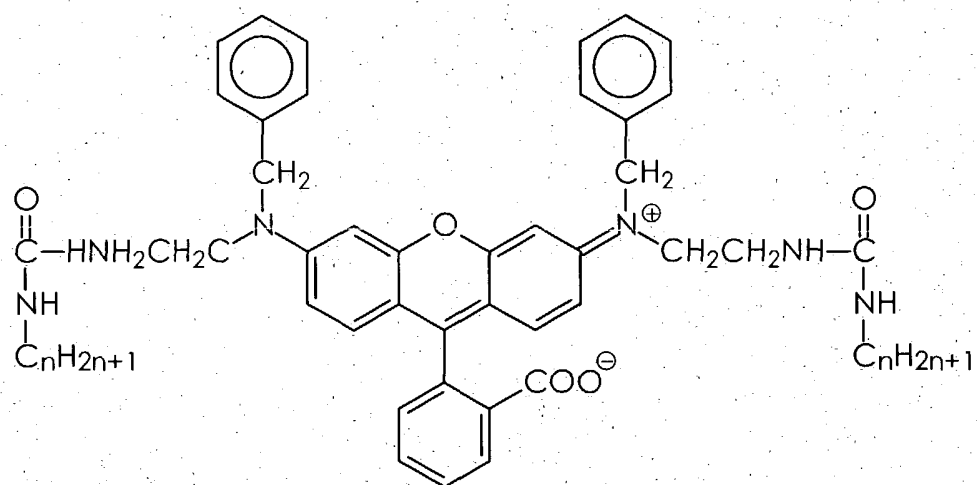
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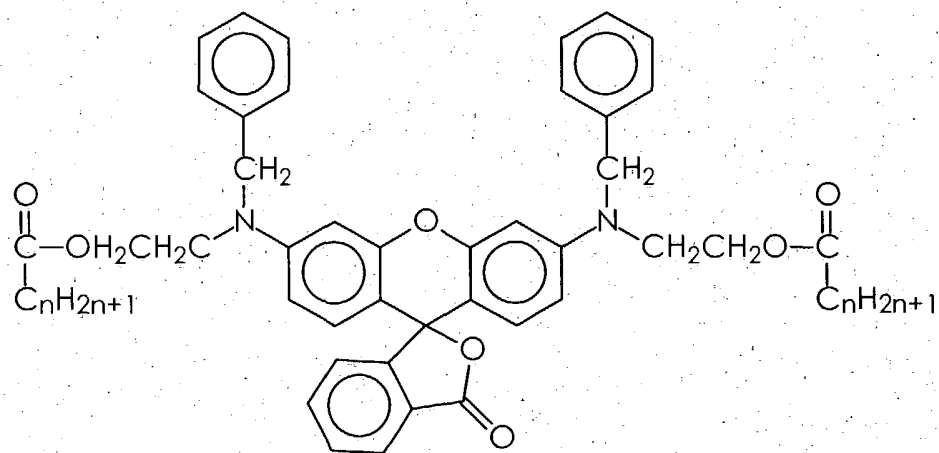
wherein n is at least about 12, (I) those of the formulae

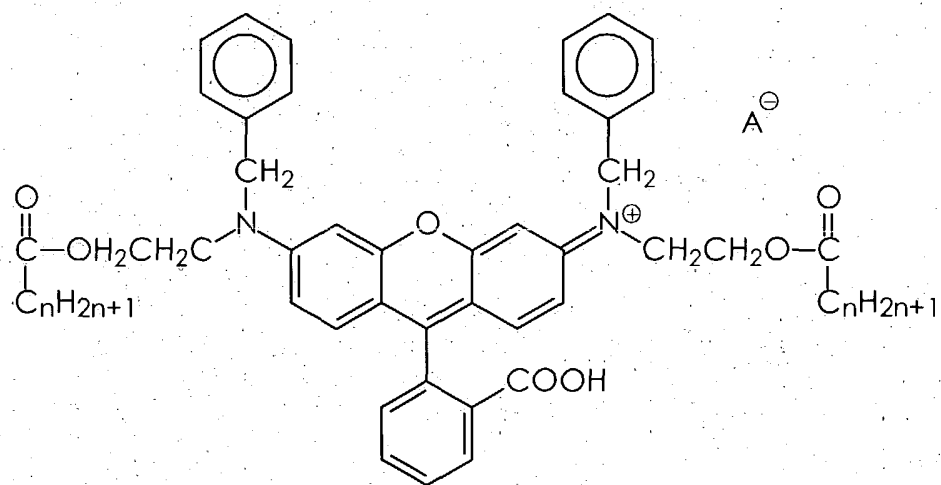


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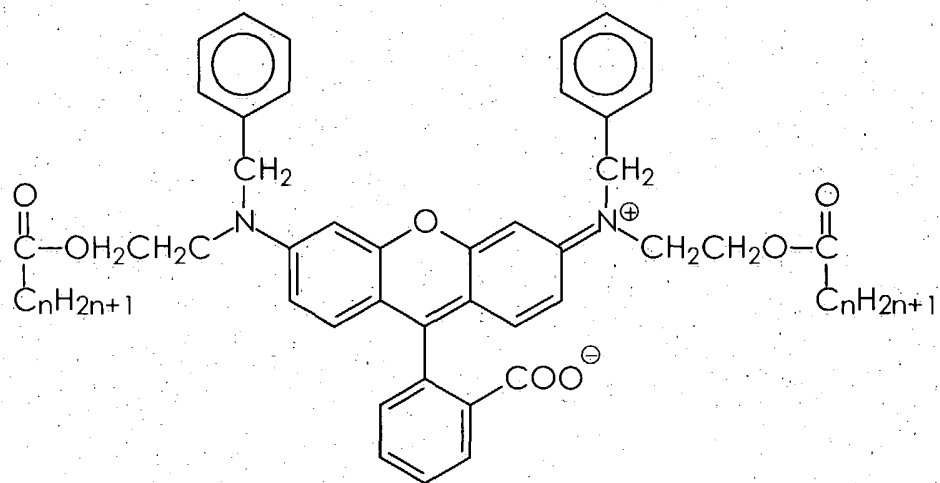


wherein n is at least about 12, (m) those of the formulae

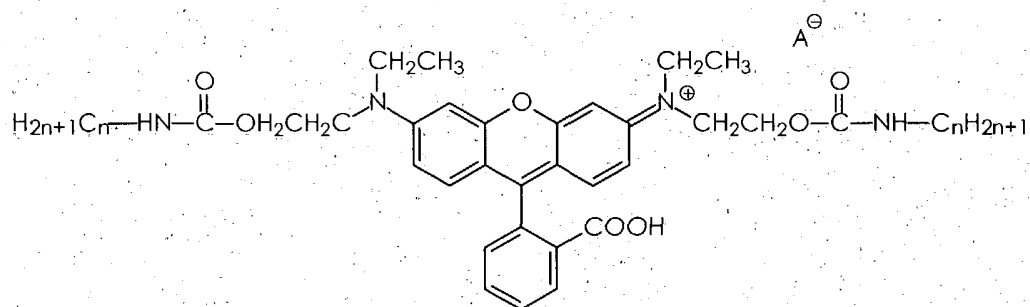
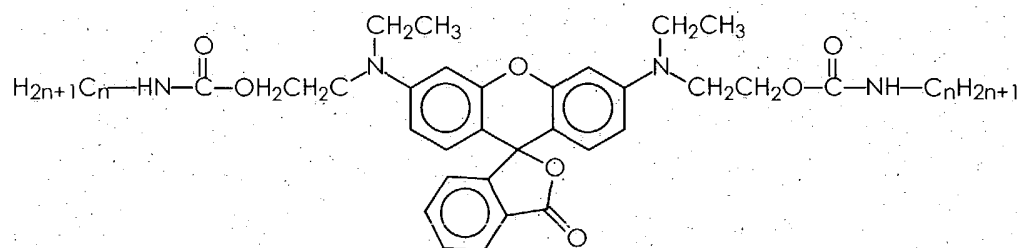




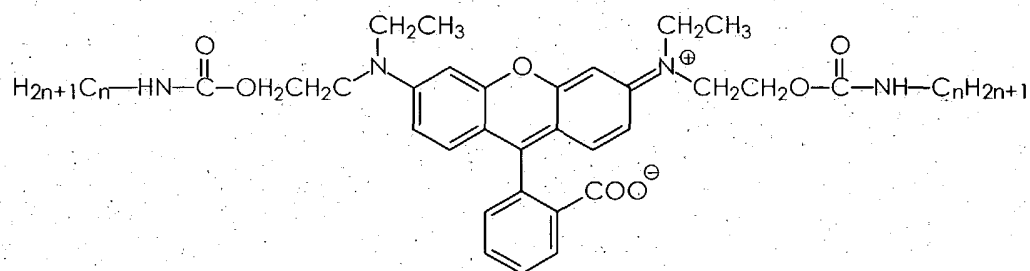
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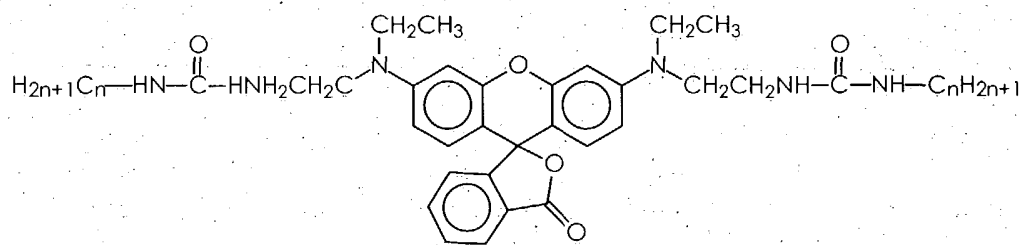
wherein n is at least about 12, (n) those of the formulae

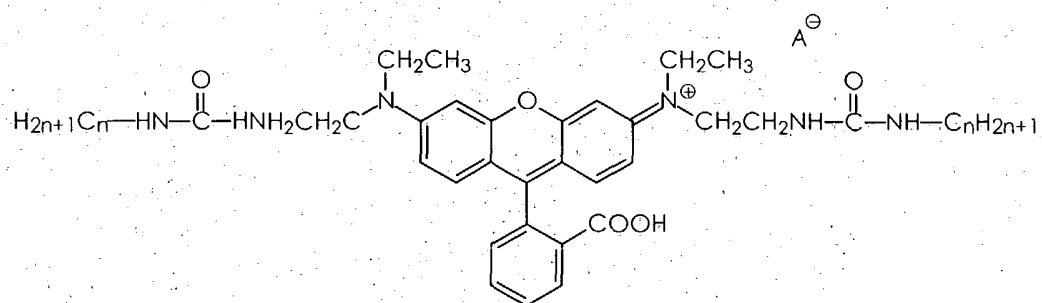


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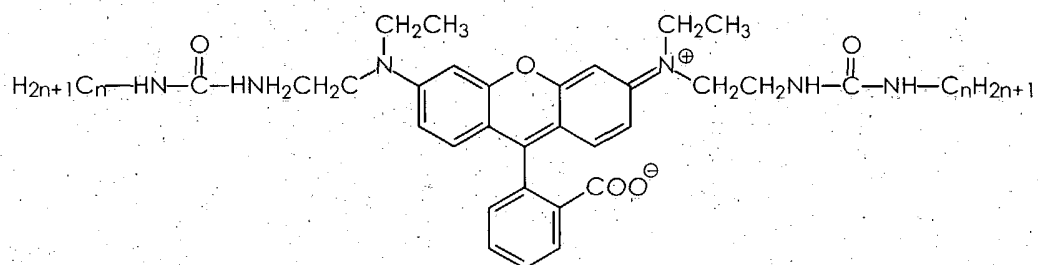


5 wherein n is at least about 12, (o) those of the formulae

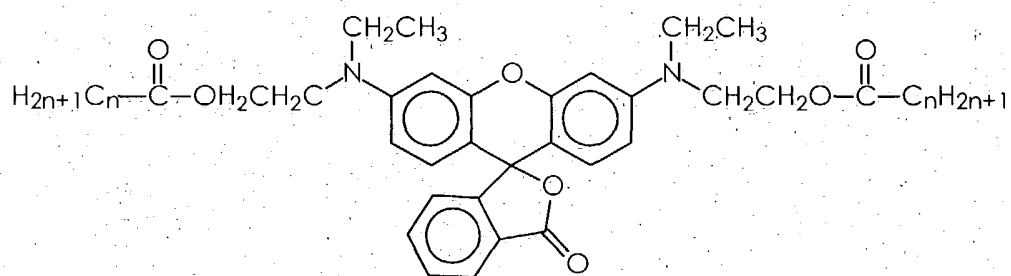


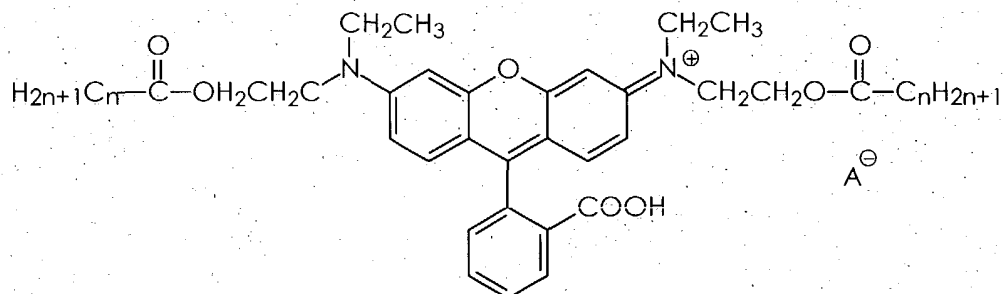


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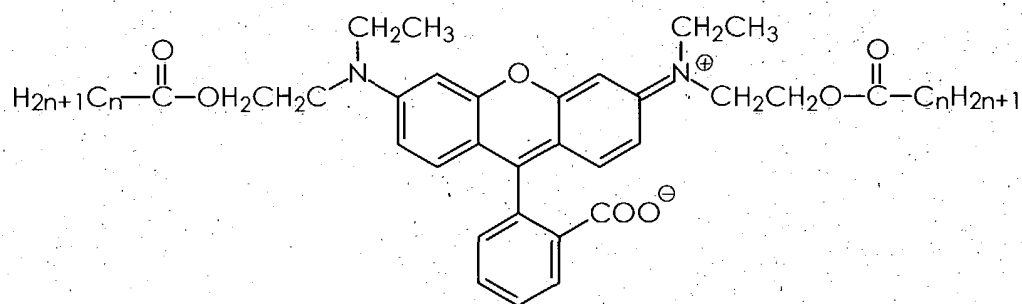


wherein n is at least about 12, (p) those of the formulae





and



wherein n is at least about 12, and the like.

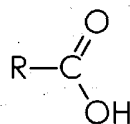
5 Phase change inks of the present invention contain a phase change carrier system or composition. The phase change carrier composition is typically designed for use in either a direct printing mode or an indirect or offset printing transfer system.

In the direct printing mode, the phase change carrier
 10 composition in one embodiment contains one or more materials that enable the phase change ink (1) to be applied in a thin film of uniform thickness on the final recording substrate (such as paper, transparency material, and the like) when cooled to ambient temperature after printing directly to the recording substrate, (2) to be ductile while
 15 retaining sufficient flexibility so that the applied image on the substrate

will not fracture upon bending, and (3) to possess a high degree of lightness, chroma, transparency, and thermal stability.

In an offset printing transfer or indirect printing mode, the phase change carrier composition in one embodiment exhibits not only the characteristics desirable for direct printing mode inks, but also certain fluidic and mechanical properties desirable for use in such a system, as described in, for example, U.S. Patent 5,389,958 the disclosure of which is totally incorporated herein by reference.

Any desired or effective carrier composition can be used. Examples of suitable ink carrier materials include fatty amides, such as monoamides, tetra-amides, mixtures thereof, and the like. Specific examples of suitable fatty amide ink carrier materials include stearyl stearamide, a dimer acid based tetra-amide that is the reaction product of dimer acid, ethylene diamine, and stearic acid, a dimer acid based tetra-amide that is the reaction product of dimer acid, ethylene diamine, and a carboxylic acid having at least about 36 carbon atoms, and the like, as well as mixtures thereof. When the fatty amide ink carrier is a dimer acid based tetra-amide that is the reaction product of dimer acid, ethylene diamine, and a carboxylic acid having at least about 36 carbon atoms, the carboxylic acid is of the general formula



wherein R is an alkyl group, including linear, branched, saturated, unsaturated, and cyclic alkyl groups, said alkyl group in one embodiment having at least about 36 carbon atoms, in another

embodiment having at least about 40 carbon atoms, said alkyl group in one embodiment having no more than about 200 carbon atoms, in another embodiment having no more than about 150 carbon atoms, and in yet another embodiment having no more than about 100 carbon atoms, although the number of carbon atoms can be outside of these ranges. Carboxylic acids of this formula are commercially available from, for example, Baker Petrolite, Tulsa, OK, and can also be prepared as described in Example 1 of U.S. Patent 6,174,937, the disclosure of which is totally incorporated herein by reference. Further information on fatty amide carrier materials is disclosed in, for example, U.S. Patent 4,889,560, U.S. Patent 4,889,761, U.S. Patent 5,194,638, U.S. Patent 4,830,671, U.S. Patent 6,174,937, U.S. Patent 5,372,852, U.S. Patent 5,597,856, U.S. Patent 6,174,937, and British Patent GB 2 238 792, the disclosures of each of which are totally incorporated herein by reference.

Also suitable as phase change ink carrier materials are isocyanate-derived resins and waxes, such as urethane isocyanate-derived materials, urea isocyanate-derived materials, urethane/urea isocyanate-derived materials, mixtures thereof, and the like. Further information on isocyanate-derived carrier materials is disclosed in, for example, U.S. Patent 5,750,604, U.S. Patent 5,780,528, U.S. Patent 5,782,966, U.S. Patent 5,783,658, U.S. Patent 5,827,918, U.S. Patent 5,830,942, U.S. Patent 5,919,839, U.S. Patent 6,255,432, U.S. Patent 6,309,453, British Patent GB 2 294 939, British Patent GB 2 305 928, British Patent GB 2 305 670, British Patent GB 2 290 793, PCT Publication WO 94/14902, PCT Publication WO 97/12003, PCT Publication WO 97/13816, PCT Publication WO 96/14364, PCT Publication

WO 97/33943, and PCT Publication WO 95/04760, the disclosures of each of which are totally incorporated herein by reference.

Mixtures of fatty amide materials and isocyanate-derived materials can also be employed as the ink carrier composition for inks
5 of the present invention.

Additional suitable phase change ink carrier materials for the present invention include paraffins, microcrystalline waxes, polyethylene waxes, ester waxes, amide waxes, fatty acids, fatty alcohols, fatty amides and other waxy materials, sulfonamide materials,
10 resinous materials made from different natural sources (such as, for example, tall oil rosins and rosin esters), and many synthetic resins, oligomers, polymers and copolymers, such as ethylene/vinyl acetate copolymers, ethylene/acrylic acid copolymers, ethylene/vinyl acetate/acrylic acid copolymers, copolymers of acrylic acid with
15 polyamides, and the like, ionomers, and the like, as well as mixtures thereof. One or more of these materials can also be employed in a mixture with a fatty amide material and/or an isocyanate-derived material.

In one specific embodiment, the phase change ink carrier
20 comprises (a) a polyethylene wax, present in the ink in an amount in one embodiment of at least about 25 percent by weight of the ink, in another embodiment of at least about 30 percent by weight of the ink, and in yet another embodiment of at least about 37 percent by weight of the ink, and in one embodiment of no more than about 60 percent
25 by weight of the ink, in another embodiment of no more than about 53 percent by weight of the ink, and in yet another embodiment of no more than about 48 percent by weight of the ink, although the amount

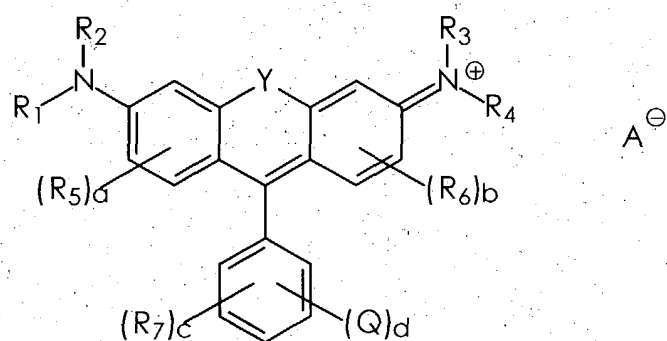
can be outside of these ranges; (b) a stearyl stearamide wax, present in the ink in an amount in one embodiment of at least about 8 percent by weight of the ink, in another embodiment of at least about 10 percent by weight of the ink, and in yet another embodiment of at least about 12 percent by weight of the ink, and in one embodiment of no more than about 32 percent by weight of the ink, in another embodiment of no more than about 28 percent by weight of the ink, and in yet another embodiment of no more than about 25 percent by weight of the ink, although the amount can be outside of these ranges; (c) a dimer acid based tetra-amide that is the reaction product of dimer acid, ethylene diamine, and a long chain hydrocarbon having greater than thirty six carbon atoms and having a terminal carboxylic acid group, present in the ink in an amount in one embodiment of at least about 10 percent by weight of the ink, in another embodiment of at least about 13 percent by weight of the ink, and in yet another embodiment of at least about 16 percent by weight of the ink, and in one embodiment of no more than about 32 percent by weight of the ink, in another embodiment of no more than about 27 percent by weight of the ink, and in yet another embodiment of no more than about 22 percent by weight of the ink, although the amount can be outside of these ranges; (d) a urethane resin derived from the reaction of two equivalents of hydroabietyl alcohol and one equivalent of isophorone diisocyanate, present in the ink in an amount in one embodiment of at least about 6 percent by weight of the ink, in another embodiment of at least about 8 percent by weight of the ink, and in yet another embodiment of at least about 10 percent by weight of the ink, and in one embodiment of no more than about 16 percent by weight of the ink, in another

embodiment of no more than about 14 percent by weight of the ink, and in yet another embodiment of no more than about 12 percent by weight of the ink, although the amount can be outside of these ranges; (e) a urethane resin that is the adduct of three equivalents of stearyl isocyanate and a glycerol-based propoxylate alcohol, present in the ink in an amount in one embodiment of at least about 2 percent by weight of the ink, in another embodiment of at least about 3 percent by weight of the ink, and in yet another embodiment of at least about 4.5 percent by weight of the ink, and in one embodiment of no more than about 13 percent by weight of the ink, in another embodiment of no more than about 10 percent by weight of the ink, and in yet another embodiment of no more than about 7.5 percent by weight of the ink, although the amount can be outside of these ranges; and (f) an antioxidant, present in the ink in an amount in one embodiment of at least about 0.01 percent by weight of the ink, in another embodiment of at least about 0.05 percent by weight of the ink, and in yet another embodiment of at least about 0.1 percent by weight of the ink, and in one embodiment of no more than about 1 percent by weight of the ink, in another embodiment of no more than about 0.5 percent by weight of the ink, and in yet another embodiment of no more than about 0.3 percent by weight of the ink, although the amount can be outside of these ranges.

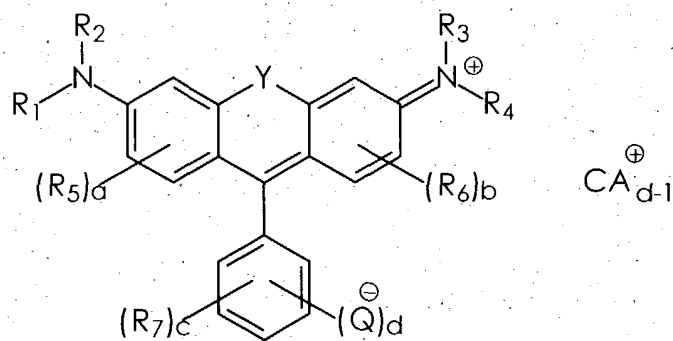
The ink carrier is present in the phase change ink of the present invention in any desired or effective amount, in one embodiment of at least about 0.1 percent by weight of the ink, in another embodiment of at least about 50 percent by weight of the ink, and in yet another embodiment of at least about 90 percent by weight

of the ink, and in one embodiment of no more than about 99 percent by weight of the ink, in another embodiment of no more than about 98 percent by weight of the ink, and in yet another embodiment of no more than about 95 percent by weight of the ink, although the amount
 5 can be outside of these ranges.

The phase change inks of the present invention contain a colorant compound of the formula



or



10

This colorant is present in the ink in any desired or effective amount to obtain the desired color or hue, in one embodiment of at least about 0.1 percent by weight of the ink, in another embodiment of at least about 0.5 percent by weight of the ink, in yet another embodiment of

at least about 1 percent by weight of the ink, in still another embodiment of at least about 2 percent by weight of the ink, and in another embodiment of at least about 3 percent by weight of the ink, and in one embodiment of no more than about 20 percent by weight of the ink, in another embodiment of no more than about 13 percent by weight of the ink, and in yet another embodiment of no more than about 6 percent by weight of the ink, although the amount can be outside of these ranges. The colorant according to the present invention can either be the sole colorant in the ink or can be present in combination with other colorants, such as dyes, pigments, mixtures thereof, and the like.

In a specific embodiment, the inks of the present invention include an anthraquinone colorant in addition to the colorant according to the present invention. Examples of suitable anthraquinone colorants include Solvent Red 172, colorants as disclosed in U.S. Patent 6,395,078 and U.S. Patent 6,422,695, the disclosures of each of which are totally incorporated herein by reference, colorants as disclosed in Copending Application U.S. Serial No. 10/260,146, Copending Application U.S. Serial No. 10/260,376, and Copending Application U.S. Serial No. 10/260,379, the disclosures of each of which are totally incorporated herein by reference, and the like. In a specific embodiment, the anthraquinone colorant is one prepared as described in Example XVII, Parts 1 through 5. The anthraquinone colorant can be present in the inks of the present invention in any desired or effective amount to achieve the desired color, hue, and other characteristics, in one embodiment of at least about 1 percent by weight of the ink, in another embodiment of at

least about 2 percent by weight of the ink, and in yet another embodiment of at least about 3 percent by weight of the ink, and in one embodiment of no more than about 20 percent by weight of the ink, in another embodiment of no more than about 13 percent by weight of the ink, and in yet another embodiment of no more than about 6 percent by weight of the ink, although the amount can be outside of these ranges.

In specific embodiments, the inks of the present invention further contain an acid having a K_a value greater than that of the K_a of the carboxylic acid and/or sulfonic acid and/or carboxylate and/or sulfonate groups on the colorant. Specific examples of suitable acids include organic sulfonic acids, including alkyl benzene sulfonic acids such as para-toluene-sulfonic acid, dodecylbenzenesulfonic acid, and the like, p-toluene sulfonic acid, hydrochloric acid, trifluoroacetic acid, methylsulfonic acid, trifluoromethyl sulfonic acid, hydrobromic acid, and the like, as well as mixtures thereof. The acid is present in any desired or effective amount, in one embodiment at least about 2 percent by weight of the amount of colorant according to the present invention, and in another embodiment at least about 5 percent by weight of the amount of colorant according to the present invention, and in one embodiment no more than about 100 percent by weight of the amount of the colorant according to the present invention, and in another embodiment no more than about 30 percent by weight of the colorant according to the present invention, although the amount of acid can be outside of these ranges.

The inks of the present invention can also optionally contain an antioxidant. The optional antioxidants of the ink compositions

protect the images from oxidation and also protect the ink components from oxidation during the heating portion of the ink preparation process. Specific examples of suitable antioxidants include NAUGUARD® 524, NAUGUARD® 76, and NAUGUARD® 512
5 (commercially available from Uniroyal Chemical Company, Oxford, CT), IRGANOX® 1010 (commercially available from Ciba Geigy), and the like. When present, the optional antioxidant is present in the ink in any desired or effective amount, in one embodiment of at least about 0.01 percent by weight of the ink, in another embodiment of at least
10 about 0.1 percent by weight of the ink, and in yet another embodiment of at least about 1 percent by weight of the ink, and in one embodiment of no more than about 20 percent by weight of the ink, in another embodiment of no more than about 5 percent by weight of the ink, and in yet another embodiment of no more than about 3
15 percent by weight of the ink, although the amount can be outside of these ranges.

The inks of the present invention can also optionally contain a viscosity modifier. Examples of suitable viscosity modifiers include aliphatic ketones, such as stearone, and the like. When present, the
20 optional viscosity modifier is present in the ink in any desired or effective amount, in one embodiment of at least about 0.1 percent by weight of the ink; in another embodiment of at least about 1 percent by weight of the ink, and in yet another embodiment of at least about 10 percent by weight of the ink, and in one embodiment of no more than about 99
25 percent by weight of the ink, in another embodiment of no more than about 30 percent by weight of the ink, and in yet another embodiment of no more than about 15 percent by weight of the ink, although the

amount can be outside of these ranges.

Other optional additives to the inks include clarifiers, such as UNION CAMP® X37-523-235 (commercially available from Union Camp), in an amount in one embodiment of at least about 0.01 percent by weight of the ink, in another embodiment of at least about 0.1 percent by weight of the ink, and in yet another embodiment of at least about 5 percent by weight of the ink, and in one embodiment of no more than about 98 percent by weight of the ink, in another embodiment of no more than about 50 percent by weight of the ink, and in yet another embodiment of no more than about 10 percent by weight of the ink, although the amount can be outside of these ranges, tackifiers, such as FORAL® 85, a glycerol ester of hydrogenated abietic (rosin) acid (commercially available from Hercules), FORAL® 105, a pentaerythritol ester of hydroabietic (rosin) acid (commercially available from Hercules), CELLOLYN® 21, a hydroabietic (rosin) alcohol ester of phthalic acid (commercially available from Hercules), ARAKAWA KE-311 Resin, a triglyceride of hydrogenated abietic (rosin) acid (commercially available from Arakawa Chemical Industries, Ltd.), synthetic polyterpene resins such as NEVTAC® 2300, NEVTAC® 100, and NEVTAC® 80 (commercially available from Neville Chemical Company), WINGTACK® 86, a modified synthetic polyterpene resin (commercially available from Goodyear), and the like, in an amount in one embodiment of at least about 0.1 percent by weight of the ink, in another embodiment of at least about 5 percent by weight of the ink, and in yet another embodiment of at least about 10 percent by weight of the ink, and in one embodiment of no more than about 98 percent by weight of the ink, in another embodiment of no more than about 75

percent by weight of the ink, and in yet another embodiment of no more than about 50 percent by weight of the ink, although the amount can be outside of these range, adhesives, such as VERSAMID® 757, 759, or 744 (commercially available from Henkel), in an amount in one
5 embodiment of at least about 0.1 percent by weight of the ink, in another embodiment of at least about 1 percent by weight of the ink, and in yet another embodiment of at least about 5 percent by weight of the ink, and in one embodiment of no more than about 98 percent by weight of the ink, in another embodiment of no more than about 50
10 percent by weight of the ink, and in yet another embodiment of no more than about 10 percent by weight of the ink, although the amount can be outside of these ranges, plasticizers, such as UNIPLEX® 250 (commercially available from Uniplex), the phthalate ester plasticizers commercially available from Monsanto under the trade name
15 SANTICIZER®, such as dioctyl phthalate, diundecyl phthalate, alkylbenzyl phthalate (SANTICIZER® 278), triphenyl phosphate (commercially available from Monsanto), KP-140®, a tributoxyethyl phosphate (commercially available from FMC Corporation), MORFLEX® 150, a dicyclohexyl phthalate (commercially available from Morflex
20 Chemical Company Inc.), trioctyl trimellitate (commercially available from Eastman Kodak Co.), and the like, in an amount in one embodiment of at least about 0.1 percent by weight of the ink, in another embodiment of at least about 1 percent by weight of the ink, and in yet another embodiment of at least about 2 percent by weight
25 of the ink, and in one embodiment of no more than about 50 percent by weight of the ink, in another embodiment of no more than about 30 percent by weight of the ink, and in yet another embodiment of no

more than about 10 percent by weight of the ink, although the amount can be outside of these ranges, and the like.

The ink compositions of the present invention in one embodiment have melting points of no lower than about 50°C, in
5 another embodiment of no lower than about 70°C, and in yet another embodiment of no lower than about 80°C, and have melting points in one embodiment of no higher than about 160°C, in another embodiment of no higher than about 140°C, and in yet another embodiment of no higher than about 100°C, although the melting point
10 can be outside of these ranges.

The ink compositions of the present invention generally have melt viscosities at the jetting temperature (in one embodiment no lower than about 75°C, in another embodiment no lower than about 100°C, and in yet another embodiment no lower than about 120°C,
15 and in one embodiment no higher than about 180°C, and in another embodiment no higher than about 150°C, although the jetting temperature can be outside of these ranges) in one embodiment of no more than about 30 centipoise, in another embodiment of no more than about 20 centipoise, and in yet another embodiment of no more
20 than about 15 centipoise, and in one embodiment of no less than about 2 centipoise, in another embodiment of no less than about 5 centipoise, and in yet another embodiment of no less than about 7 centipoise, although the melt viscosity can be outside of these ranges.

The ink compositions of the present invention can be
25 prepared by any desired or suitable method. For example, the ink ingredients can be mixed together, followed by heating, to a temperature in one embodiment of at least about 100°C, and in one

embodiment of no more than about 140°C, although the temperature can be outside of these ranges, and stirring until a homogeneous ink composition is obtained, followed by cooling the ink to ambient temperature (typically from about 20 to about 25°C). The inks of the present invention are solid at ambient temperature. In a specific embodiment, during the formation process, the inks in their molten state are poured into molds and then allowed to cool and solidify to form ink sticks.

The inks of the present invention can be employed in apparatus for direct printing ink jet processes and in indirect (offset) printing ink jet applications. Another embodiment of the present invention is directed to a process which comprises incorporating an ink of the present invention into an ink jet printing apparatus, melting the ink, and causing droplets of the melted ink to be ejected in an imagewise pattern onto a recording substrate. A direct printing process is also disclosed in, for example, U.S. Patent 5,195,430, the disclosure of which is totally incorporated herein by reference. Yet another embodiment of the present invention is directed to a process which comprises incorporating an ink of the present invention into an ink jet printing apparatus, melting the ink, causing droplets of the melted ink to be ejected in an imagewise pattern onto an intermediate transfer member, and transferring the ink in the imagewise pattern from the intermediate transfer member to a final recording substrate. In a specific embodiment, the intermediate transfer member is heated to a temperature above that of the final recording sheet and below that of the melted ink in the printing apparatus. An offset or indirect printing process is also disclosed in, for example, U.S. Patent 5,389,958, the

disclosure of which is totally incorporated herein by reference. In one specific embodiment, the printing apparatus employs a piezoelectric printing process wherein droplets of the ink are caused to be ejected in imagewise pattern by oscillations of piezoelectric vibrating elements.

5 Inks of the present invention can also be employed in other hot melt printing processes, such as hot melt acoustic ink jet printing, hot melt thermal ink jet printing, hot melt continuous stream or deflection ink jet printing, and the like. Phase change inks of the present invention can also be used in printing processes other than hot melt ink jet printing
10 processes.

Any suitable substrate or recording sheet can be employed, including plain papers such as XEROX® 4024 papers, XEROX® Image Series papers, Courtland 4024 DP paper, ruled notebook paper, bond paper, silica coated papers such as Sharp
15 Company silica coated paper, JuJo paper, HAMMERMILL LASERPRINT® paper, and the like, transparency materials, fabrics, textile products, plastics, polymeric films, inorganic substrates such as metals and wood, and the like.

20 Specific embodiments of the invention will now be described in detail. These examples are intended to be illustrative, and the invention is not limited to the materials, conditions, or process parameters set forth in these embodiments. All parts and percentages are by weight unless otherwise indicated.

25

EXAMPLE IA

Synthesis of Dichlorofluorescein

A mixture of fluorescein (100 grams, 0.331 mole; obtained from Aldrich Chemical Co., Milwaukee, WI) and PCl_5 (128.5 grams, 0.62 mole; obtained from Aldrich Chemical Co.) in 650 milliliters of chlorobenzene was stirred and heated to 140°C in a 1 liter round bottom flask equipped with a reflux condenser. After 6 hours of heating, the reflux condenser was replaced with a distillation setup, and POCl_3 formed during the reaction as well as the chlorobenzene were distilled off. After all of the POCl_3 and chlorobenzene were removed, 300 grams of N-methyl pyrrolidinone was added and the resulting mixture was heated to 100°C with stirring until all of the crude dichlorofluorescein dissolved. The solution was then poured into a 4 liter beaker containing 1 liter of deionized water. A tan solid precipitated out and was collected on a filter and dried in a vacuum oven. The final tan solid matched the IR, NMR, and TLC of commercially available dichlorofluorescein.

Other synthetic processes can also be used. For example, a one-pot process using DMF solvent can be employed wherein the POCl_3 intermediate is not distilled off but is removed by reaction with methanol, which also precipitates the dichlorofluorescein as a white solid. Methods using toluenesulfonylchloride, a less reactive and corrosive chlorinating agent than PCl_5 , can also be used.

EXAMPLE IB

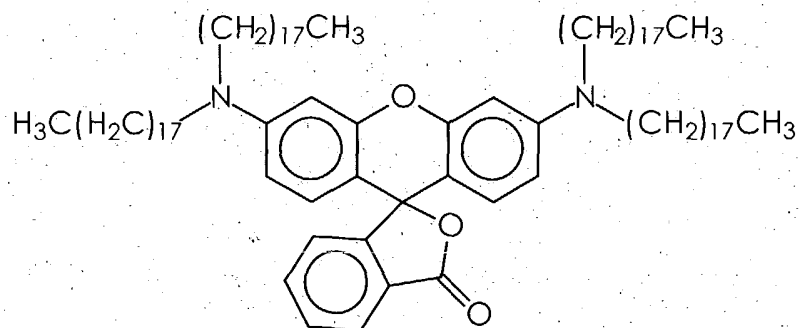
Synthesis of Tetrastearyl Colorant

A mixture of dichlorofluorescein (105 grams, 0.284 mole, prepared as described above), calcium oxide (24 grams, 0.62 mole; obtained from Aldrich Chemical Co., Milwaukee, WI), ZnCl_2 (116 grams, 0.85 mole; obtained from Aldrich Chemical Co.), and distearyl amine (288 grams, 0.585 mole; ARMEEN 2HT, obtained from Akzo-Nobel, McCook, IL) in 650 milliliters of tetramethylene sulfone (obtained from Chevron Phillips Chemical Co., LP, The Woodlands, TX) was stirred and heated to 190°C in a 1 liter round bottom flask. After 10 hours of heating, the deeply magenta colored mixture was cooled to 120°C and poured into 2.5 liters of methyl isobutyl ketone (MIBK) and stirred until totally dissolved.

EXAMPLE IC

Purification of Tetrastearyl Colorant

The solution of crude tetrastearyl colorant in MIBK was then transferred to a 4 liter separatory funnel. Three aqueous EDTA washes were then performed (50 grams of the tetrasodium salt of EDTA in 1,000 milliliters of water for each wash) to remove all of the zinc and calcium salts in the crude reaction product. The product, dissolved in MIBK, remained on the top layer with the water/EDTA chelated metal waste on the bottom layer, which was discarded. Two washes with deionized water (1 liter each) were then performed. At this point, the MIBK solution was no longer magenta, but a faint orangeish-red color. The lack of a brilliant magenta color at this point indicated a ring-closed, or free base, form of the colorant, believed to be of the formula



EXAMPLE ID

Isolation of Tetrastearyl Colorant

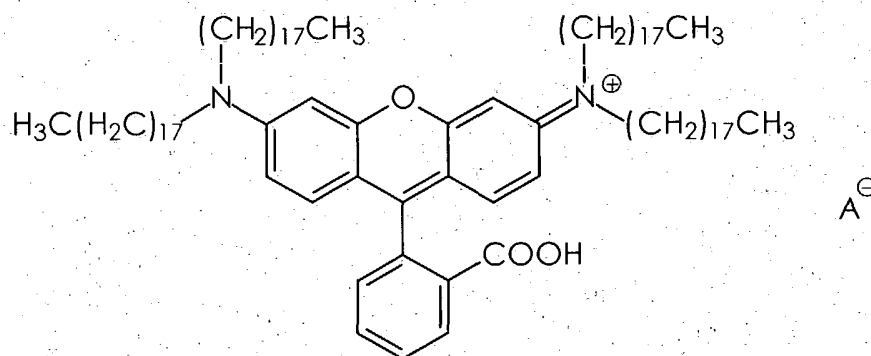
- 5 The solution of the ring-closed, purified tetrastearyl colorant in MIBK was then transferred to a 2 liter round bottom flask with distillation setup. The MIBK and residual water were distilled off and the product, a slightly viscous wax when hot, was transferred to a jar and allowed to harden. The wax was a deep red colored, somewhat hard
- 10 wax when cooled to room temperature.

EXAMPLE IE

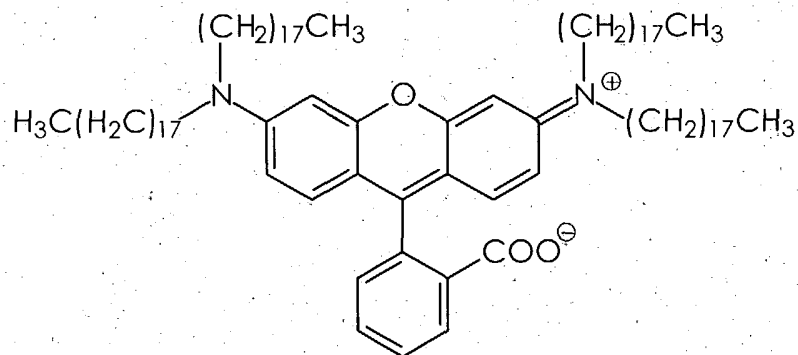
Protonation of Tetrastearyl Colorant

- 250 grams of the solid, ring-closed, purified tetrastearyl
- 15 colorant prepared in Example ID was then transferred to a 1 liter beaker and 500 milliliters of MIBK were added and allowed to dissolve the solid with stirring. A stoichiometric amount of dodecyl benzene sulfonic acid was added to this solution and stirred for 1 hour. A deep magenta hue was observed with the addition of the acid. The solution
- 20 was then transferred to a distillation setup and the MIBK removed. The molten ring-opened waxy colorant was then transferred to an

aluminum tin and allowed to cool to room temperature. The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



- 5 wherein A is the anion corresponding to the acid used for protonation. The zwitterionic form of this colorant is believed to be of the formula



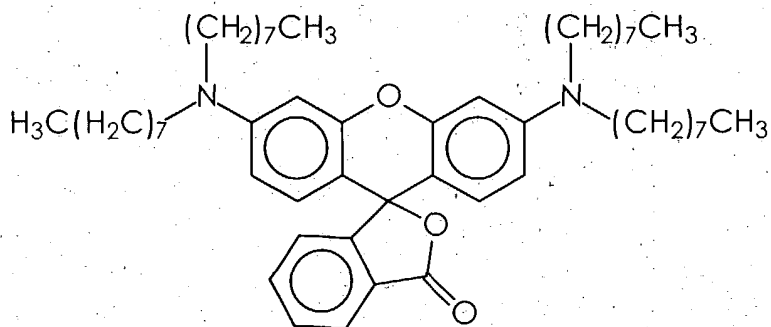
- The process was repeated a number of times substituting for dodecyl benzene sulfonic acid the following acids: p-toluene sulfonic acid; hydrochloric acid; trifluoroacetic acid; methyl sulfonic acid; trifluoromethyl sulfonic acid; and hydrobromic acid. Similar results were observed in all cases.
- 10

EXAMPLE IIB

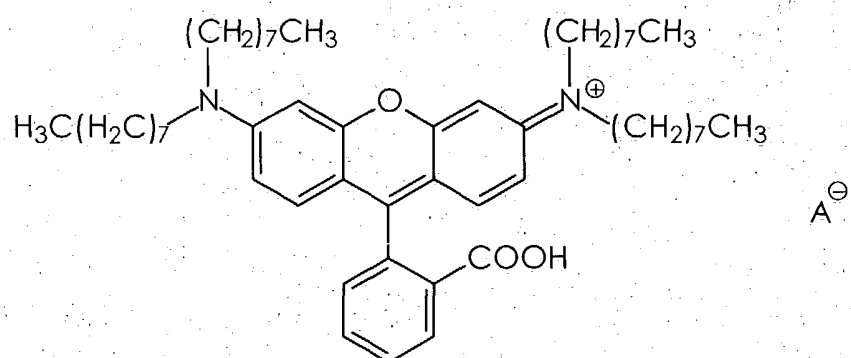
The process of Example IB was repeated except that dioctyl amine ($\text{NH}((\text{CH}_2)_7\text{CH}_3)_2$, obtained from Aldrich Chemical Co., Milwaukee, WI) was used instead of distearyl amine. The dioctyl amine was present in an amount of 1.95 moles of dioctyl amine per every one mole of dichlorofluorescein.

EXAMPLE IIC

The process of Example IC was repeated using the product obtained in Example IIB. It is believed that the purified product was of the formula

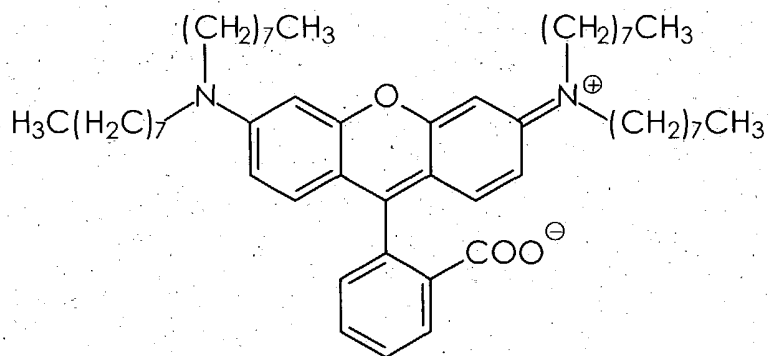


The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula



5

EXAMPLE IID

The process of Example ID was repeated using the product obtained in Example IIC.

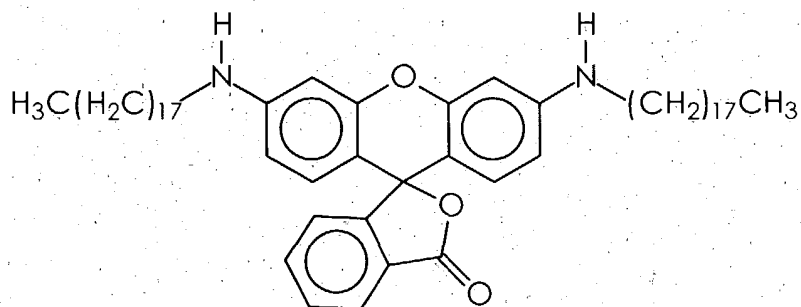
10

EXAMPLE IIIB

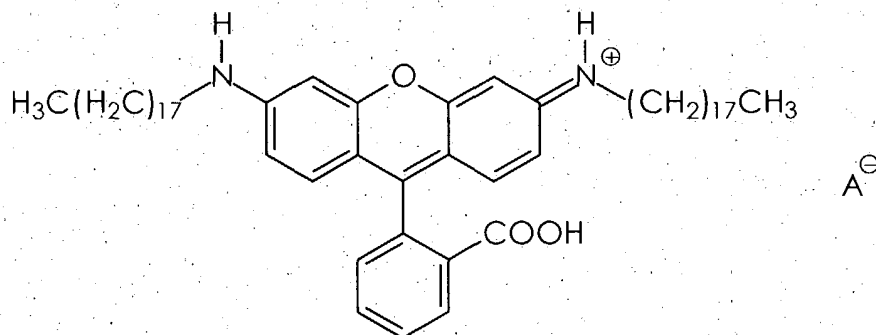
The process of Example IB was repeated except that the reaction was run with 2.05 moles of stearyl amine per every one mole of dichlorofluorescein.

EXAMPLE IIIC

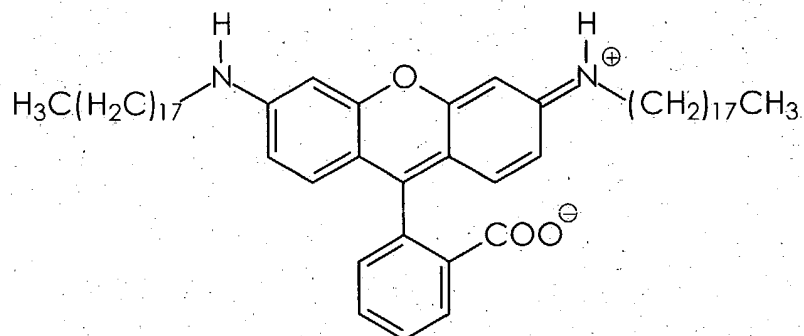
The process of Example IC was repeated using the product obtained in Example IIIB. It is believed that the purified product was of the formula



The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



10 wherein A is the anion corresponding to the acid used for protonation. The zwitterionic form of this colorant is believed to be of the formula

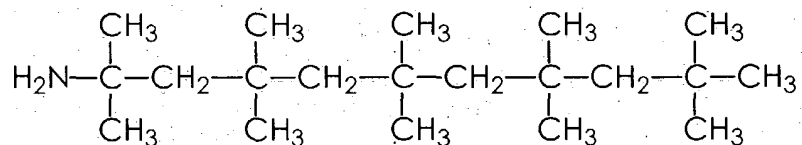


EXAMPLE IIID

The process of Example ID was repeated using the product
5 obtained in Example IIIC.

EXAMPLE IVB

The process of Example IB was repeated except that
PRIMENE JM-T (obtained from Rohm and Haas Company, Philadelphia,
10 PA), of the formula

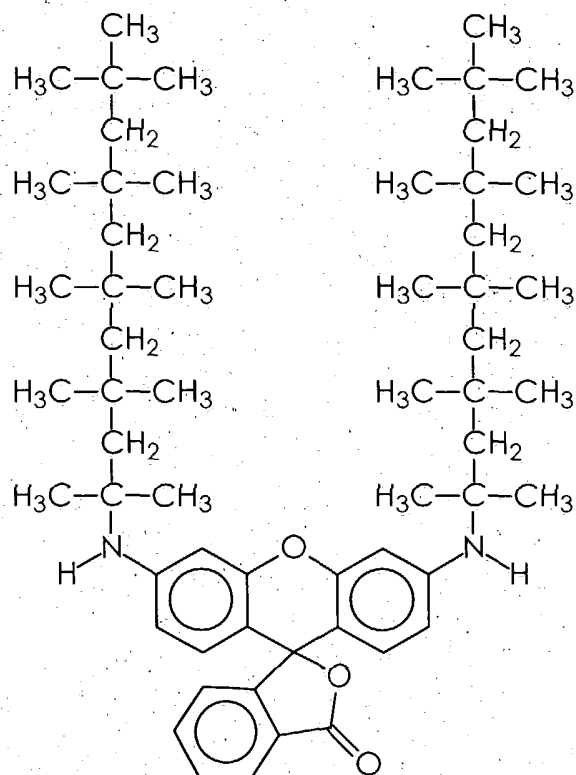


was used instead of distearyl amine. The PRIMENE JM-T was present in
an amount of 2 moles of PRIMENE JM-T per every one mole of
dichlorofluorescein.

15

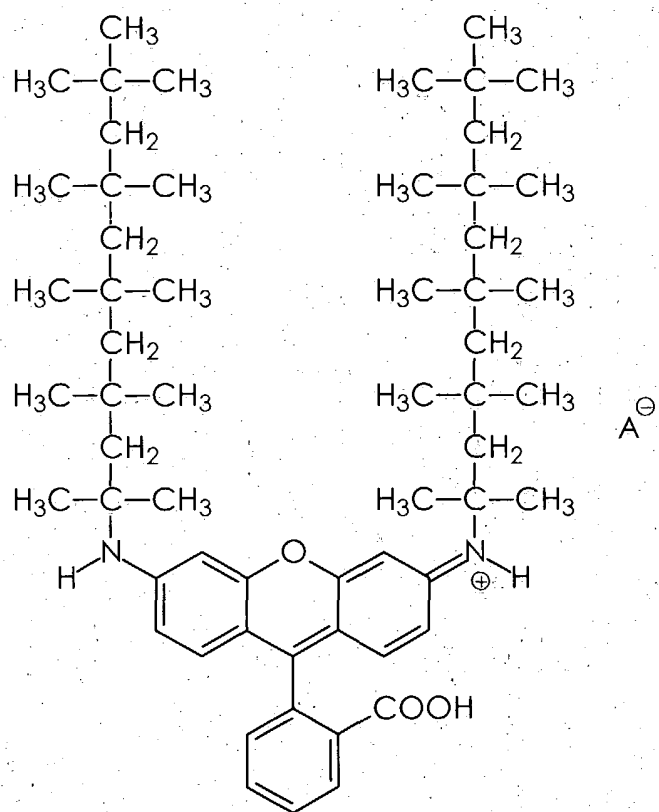
EXAMPLE IVC

The process of Example IC was repeated using the product obtained in Example IVB. It is believed that the purified product was of the formula



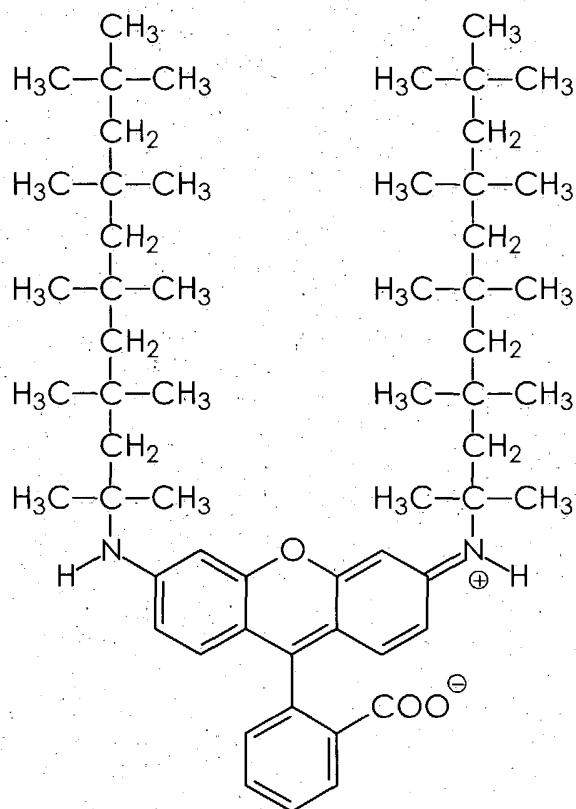
5

The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula



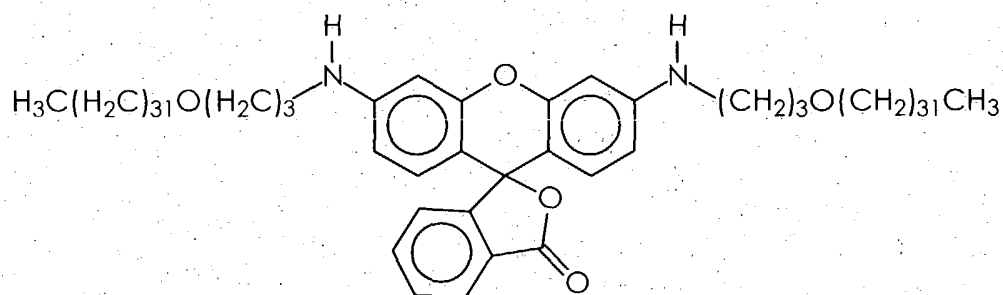
EXAMPLE IVD

The process of Example ID was repeated using the product
 5 obtained in Example IVC.

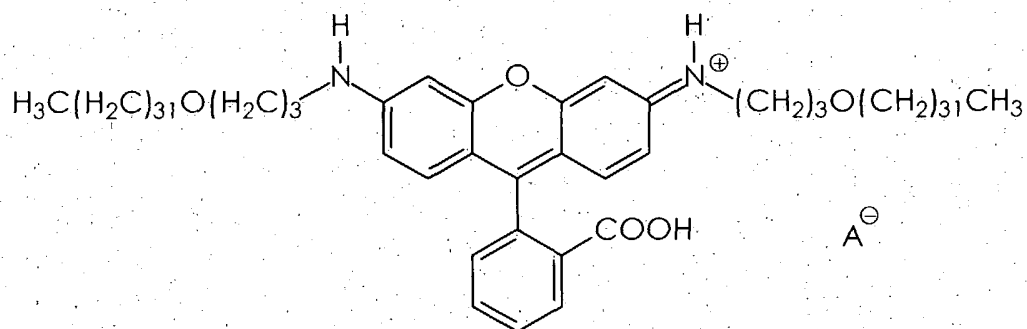
EXAMPLE VB

The process of Example IB was repeated except that
 UNILIN 425-PA (obtained from Tomah Products, Milton, WI, of the
 10 formula $\text{CH}_3(\text{CH}_2)_{31}\text{-O-CH}_2\text{CH}_2\text{CH}_2\text{NH}_2$) was used instead of distearyl
 amine. The UNILIN 425-PA was present in an amount of 2 moles of

UNILIN 425-PA per every one mole of dichlorofluorescein. It is believed that the product was of the formula

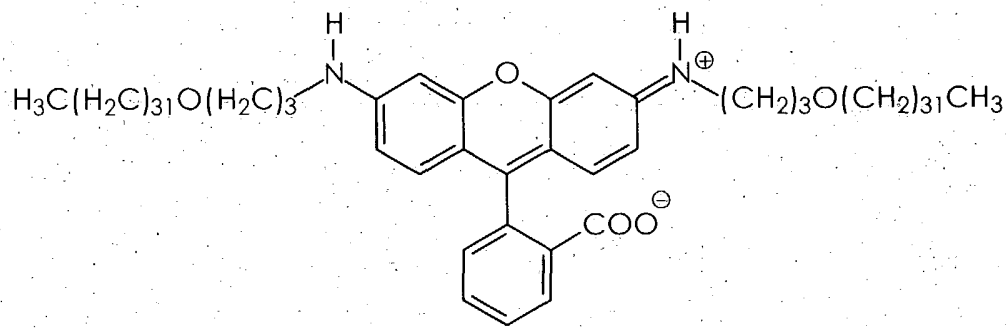


The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula

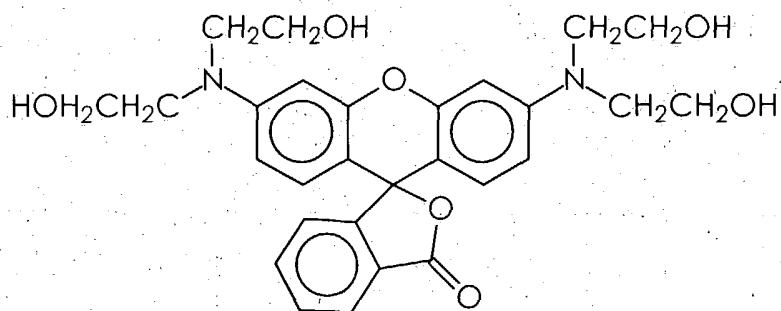


EXAMPLE VIB

The process of Example IB was repeated except that diethanol amine (obtained from Aldrich Chemical Co., Milwaukee, WI, of the formula $\text{HN}(\text{CH}_2\text{CH}_2\text{OH})_2$) was used instead of distearyl amine. The diethanol amine was present in an amount of 2.5 moles of diethanol amine per every one mole of dichlorofluorescein. In addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1 mole of calcium oxide was used per every one mole of dichlorofluorescein, the solvent was N-methyl pyrrolidone instead of tetramethylene sulfone, and the reaction mixture was heated to 125°C for 100 hours.

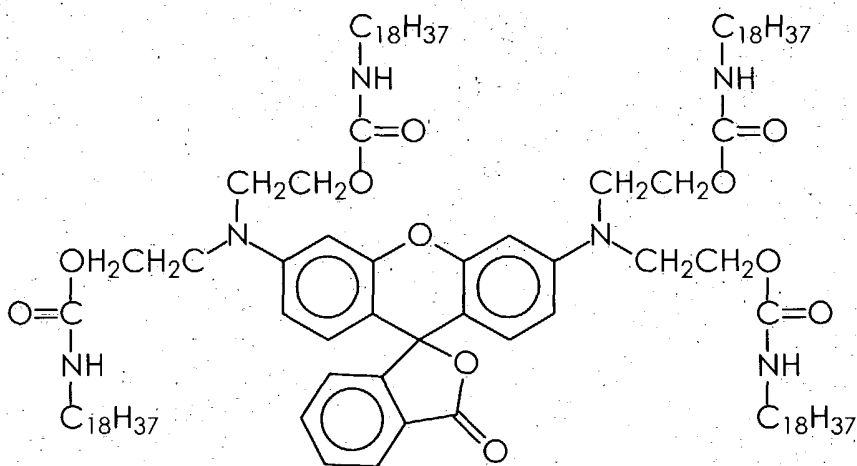
EXAMPLE VIC

The process of Example IC was repeated using the product obtained in Example VIB except that the product was poured into methanol and sufficient EDTA was added to remove all of the Zn^{2+} and Ca^{2+} ions. It is believed that the purified product was of the formula



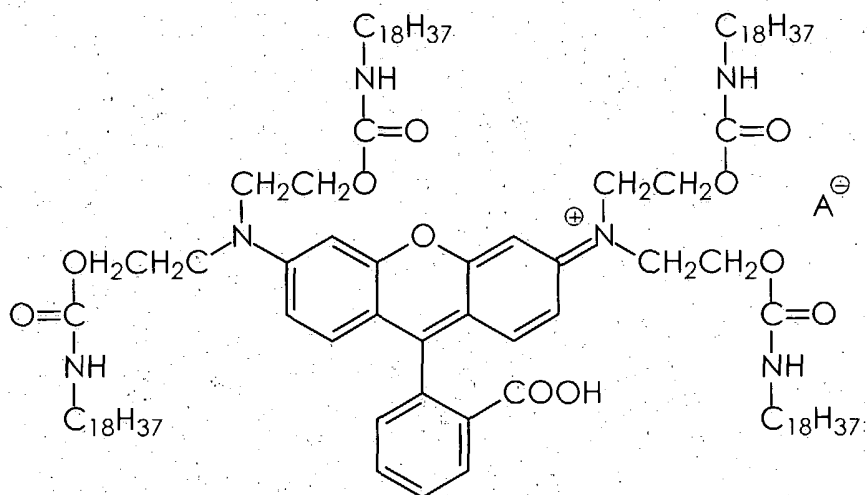
EXAMPLE VIC-1

About 10 grams of the product obtained in Example VIC is added to 23.4 grams of octadecylisocyanate (available from Aldrich Chemical Co., Milwaukee, WI) at 120°C, after which 2 drops of dibutyltindilaurate catalyst (available from Aldrich Chemical Co.) is added and the reaction is stirred and heated until disappearance of the isocyanate peak in the IR is observed. The tetraurethane rhodamine is poured into aluminum tins and is believed to be of the formula



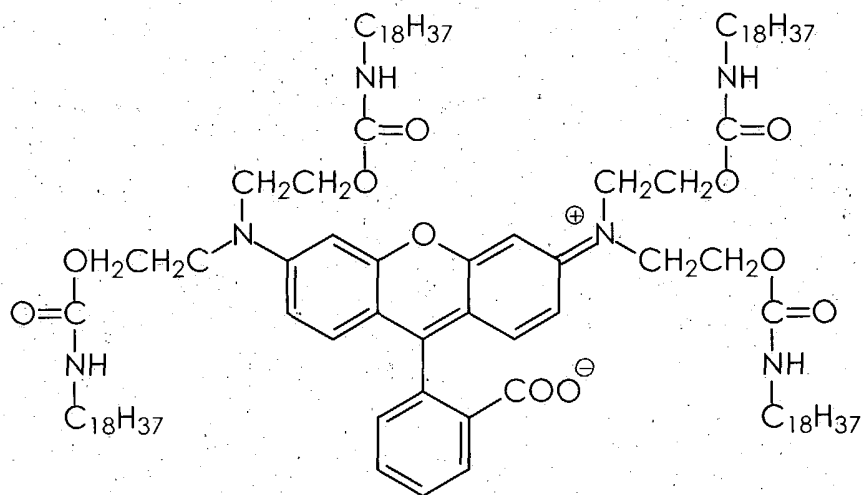
10

The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



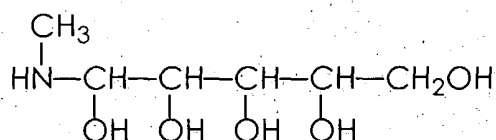
wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula



EXAMPLE VIIB

The process of Example IB was repeated except that N-methyl-D-glucamine (obtained from Aldrich Chemical Co., Milwaukee, WI), of the formula



5

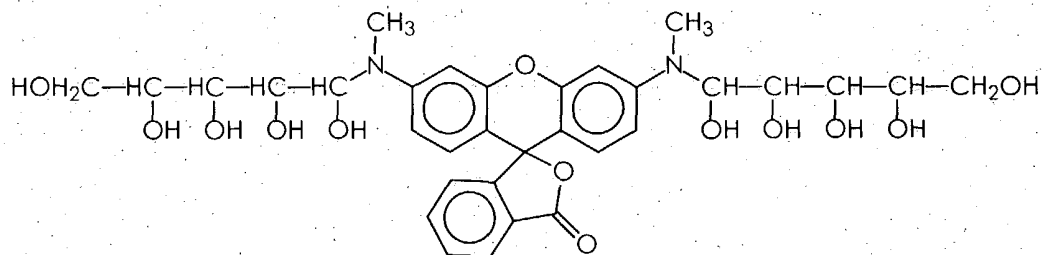
was used instead of distearyl amine. The N-methyl-D-glucamine was present in an amount of 2.5 moles of N-methyl-D-glucamine per every one mole of dichlorofluorescein. In addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1.5 moles of calcium oxide was used per every one mole of dichlorofluorescein, the solvent was N-methyl pyrrolidone instead of tetramethylene sulfone, and the reaction mixture was heated to 130°C for 7 days.

10

EXAMPLE VIIC

The process of Example IC was repeated using the product obtained in Example VIIB except that the product was poured into methanol and sufficient EDTA was added to remove all of the Zn^{2+} and Ca^{2+} ions. It is believed that the purified product was of the formula

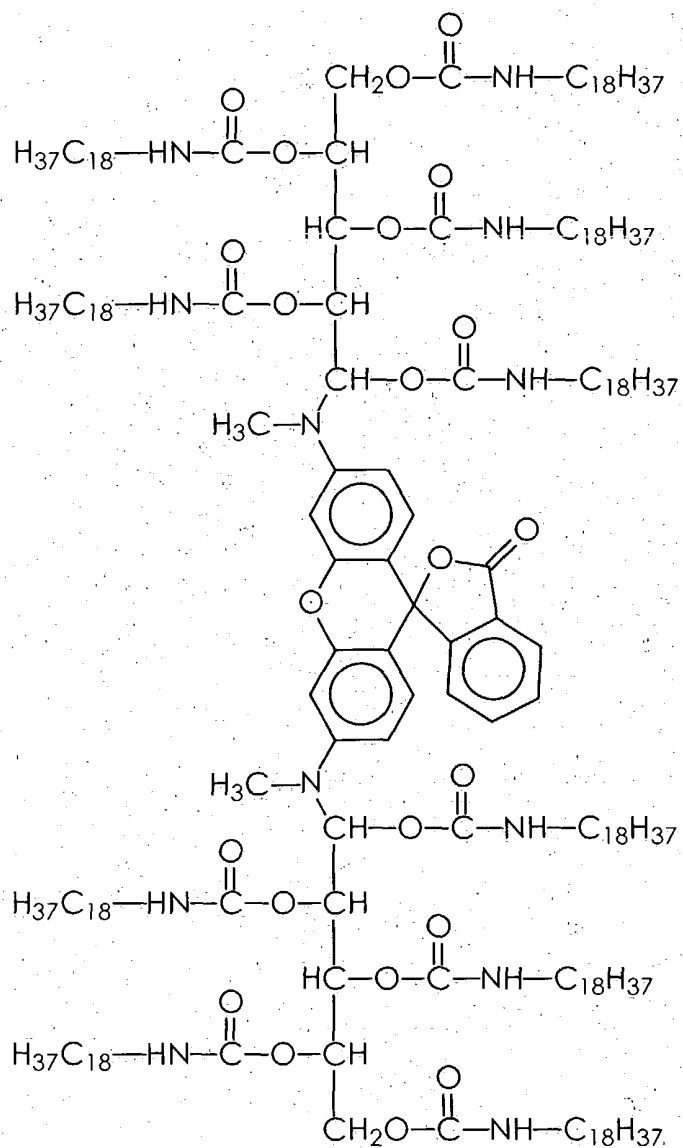
15



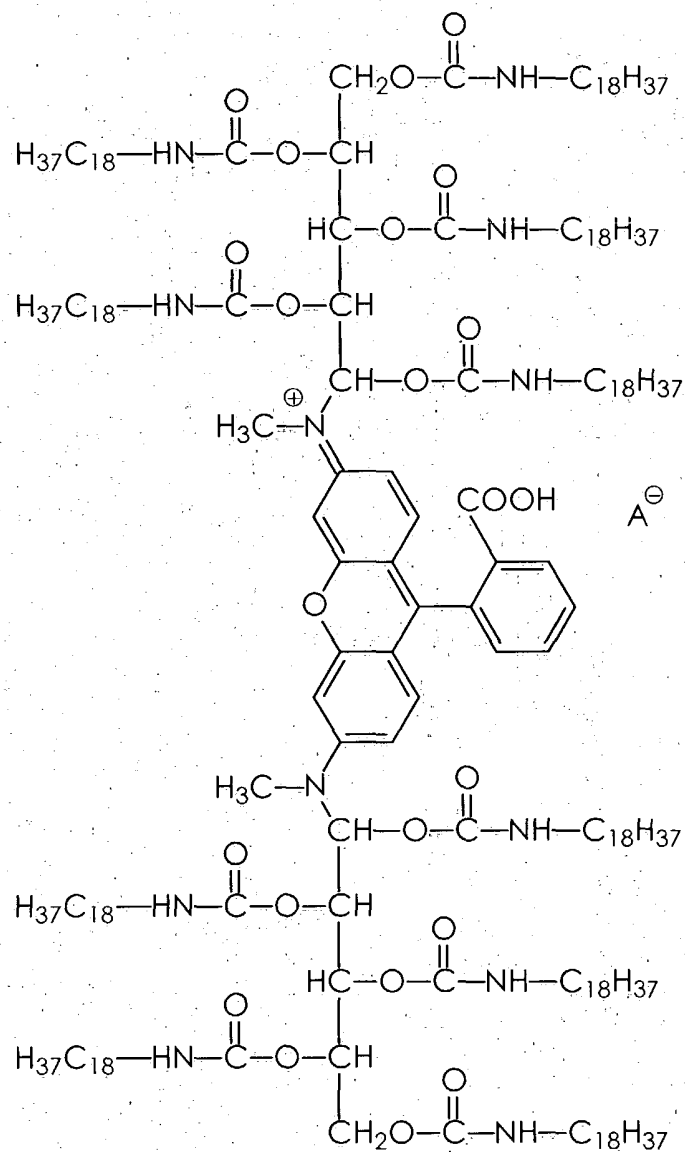
20

EXAMPLE VIIC-1

About 10 grams of the product obtained in Example VIIC is added to 45 grams of octadecylisocyanate (available from Aldrich Chemical Co., Milwaukee, WI) at 120°C, after which 4 drops of
5 dibutyltindilaurate catalyst (available from Aldrich Chemical Co.) is added and the reaction is stirred and heated until disappearance of the isocyanate peak in the IR is observed. The deca-urethane rhodamine is poured into aluminum tins and is believed to be of the formula

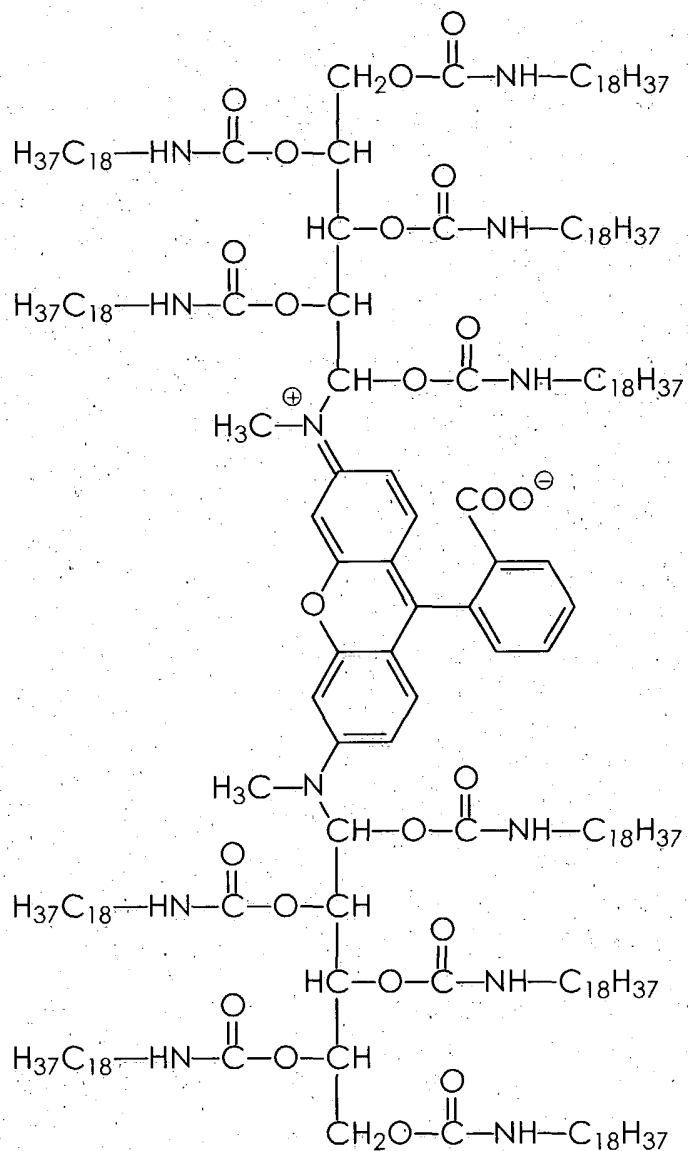


The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



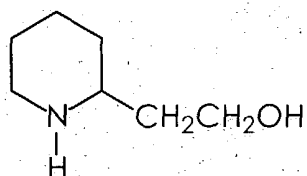
wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula

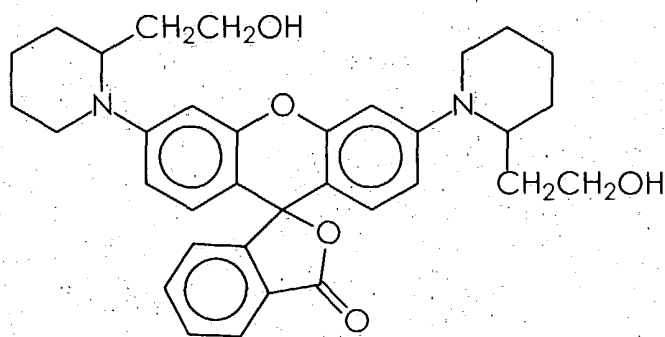


EXAMPLE VIII B

The process of Example IB was repeated except that 2-
 5 piperidine ethanol (obtained from Aldrich Chemical Co., Milwaukee,
 WI), of the formula



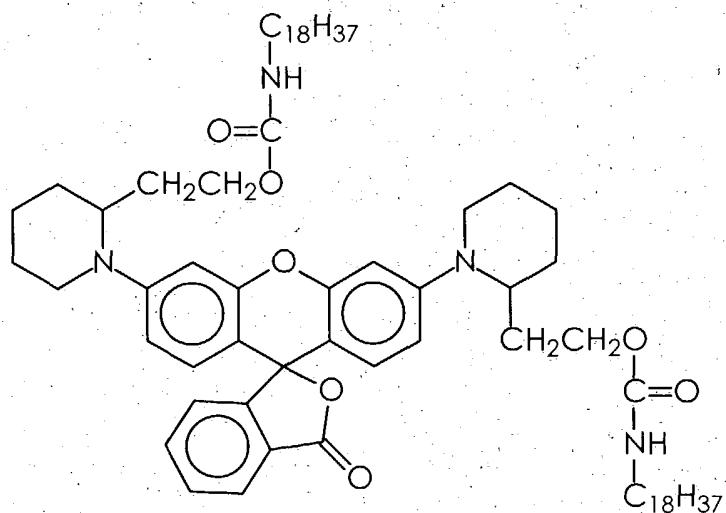
was used instead of distearyl amine. The 2-piperidine ethanol was present in an amount of 2.5 moles of 2-piperidine ethanol per every one mole of dichlorofluorescein. In addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1 mole of calcium oxide was used per every one mole of dichlorofluorescein, the solvent was N-methyl pyrrolidone instead of tetramethylene sulfone, and the reaction mixture was heated to 160°C for 24 hours. The reaction product was then poured into water and filtered and washed with water. It is believed that the product was of the formula



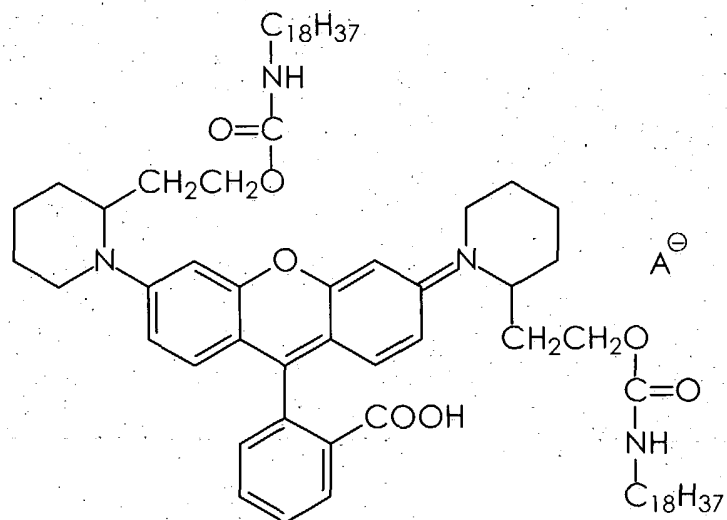
EXAMPLE VIIIC-1

About 10 grams of the product obtained in Example VIIIB is added to 10.7 grams of octadecylisocyanate (available from Aldrich Chemical Co., Milwaukee, WI) at 120°C, after which 1 drop of dibutyltindilaurate catalyst (available from Aldrich Chemical Co.) is added and the reaction is stirred and heated until disappearance of

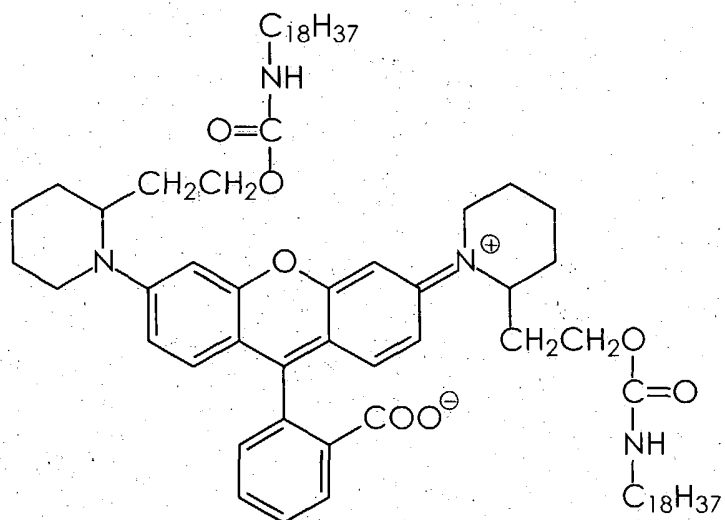
the isocyanate peak in the IR is observed. The di-urethane rhodamine is poured into aluminum tins and is believed to be of the formula



The ring-opened, or protonated, or free-base form of this colorant is
5 believed to be of the formula

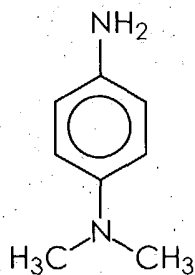


wherein A is the anion corresponding to the acid used for protonation.
The zwitterionic form of this colorant is believed to be of the formula



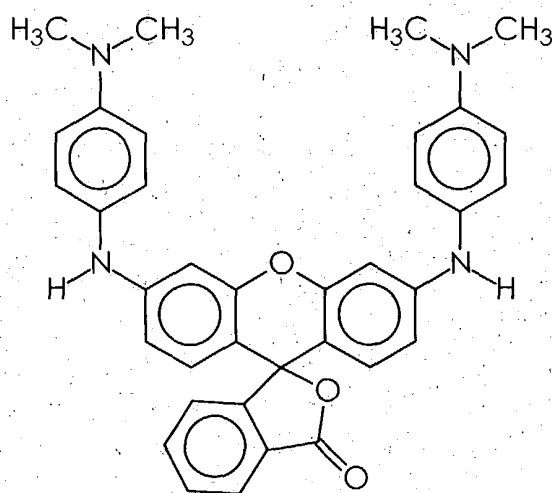
EXAMPLE IXB

The process of Example IB was repeated except that N,N-
 5 dimethyl-1,4-phenylene diamine (obtained from Aldrich Chemical Co.,
 Milwaukee, WI), of the formula



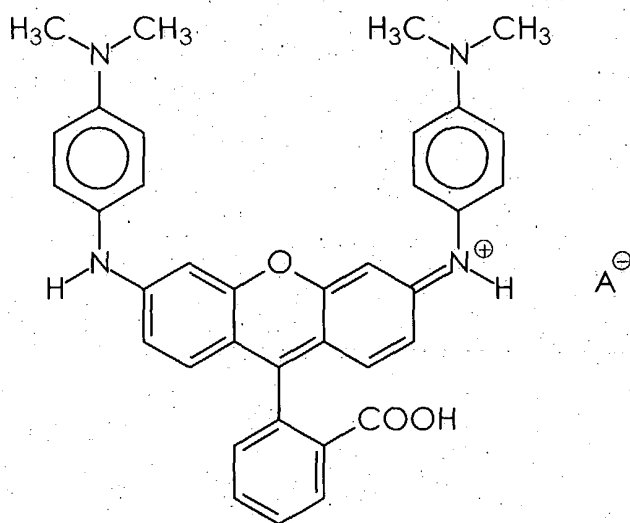
was used instead of distearyl amine. The N,N-dimethyl-1,4-phenylene
 diamine was present in an amount of 2.5 moles of N,N-dimethyl-1,4-
 10 phenylene diamine per every one mole of dichlorofluorescein. In
 addition, 2 moles of zinc chloride were used per every one mole of
 dichlorofluorescein and 1 mole of calcium oxide was used per every
 one mole of dichlorofluorescein, the solvent was N-methyl pyrrolidone

instead of tetramethylene sulfone, and the reaction mixture was heated to 140°C for 48 hours. The reaction product was then poured into water and filtered and washed with water. It is believed that the product was of the formula

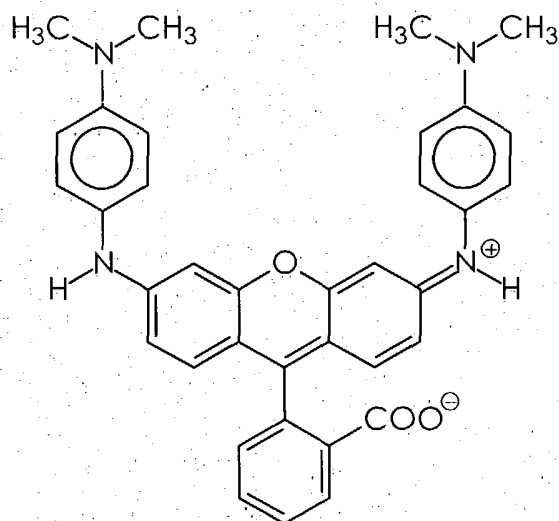


5

The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



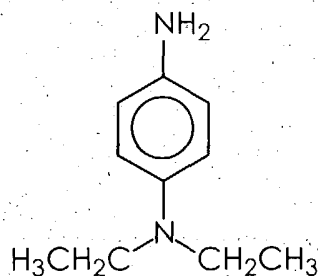
wherein A is the anion corresponding to the acid used for protonation. The zwitterionic form of this colorant is believed to be of the formula



5

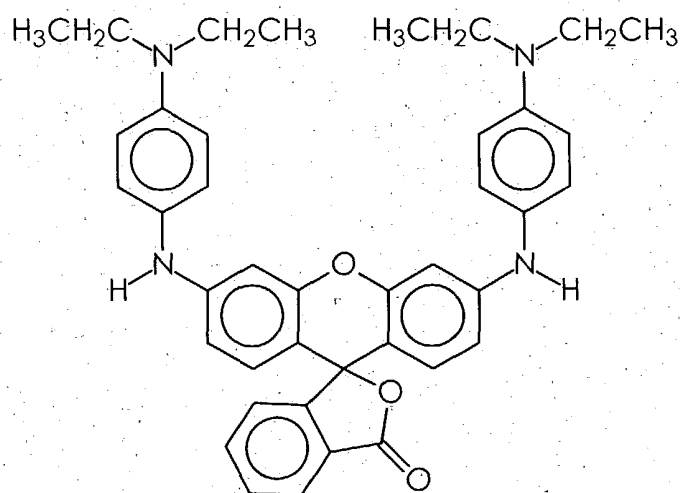
EXAMPLE XB

The process of Example IB was repeated except that N,N-diethyl-1,4-phenylene diamine (obtained from Aldrich Chemical Co., Milwaukee, WI), of the formula

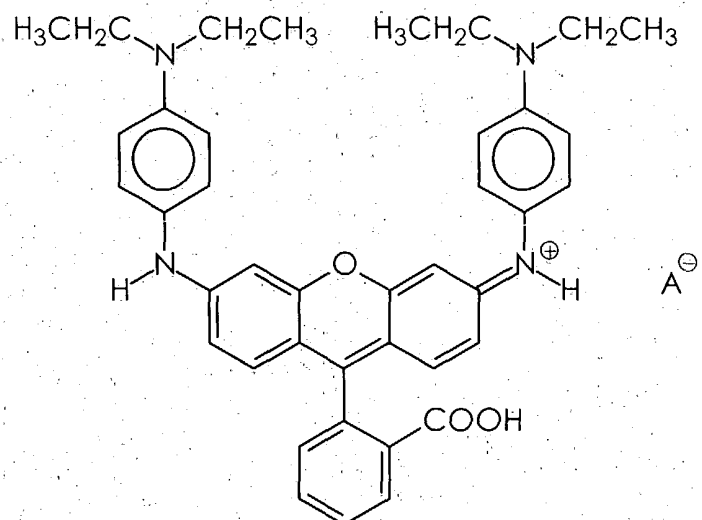


10 was used instead of distearyl amine. The N,N-diethyl-1,4-phenylene diamine was present in an amount of 2.5 moles of N,N-diethyl-1,4-phenylene diamine per every one mole of dichlorofluorescein. In

addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1 mole of calcium oxide was used per every one mole of dichlorofluorescein, the solvent was N-methyl pyrrolidone instead of tetramethylene sulfone, and the reaction mixture was
5 heated to 150°C for 96 hours. The reaction product was then poured into water and filtered and washed with water. It is believed that the product was of the formula

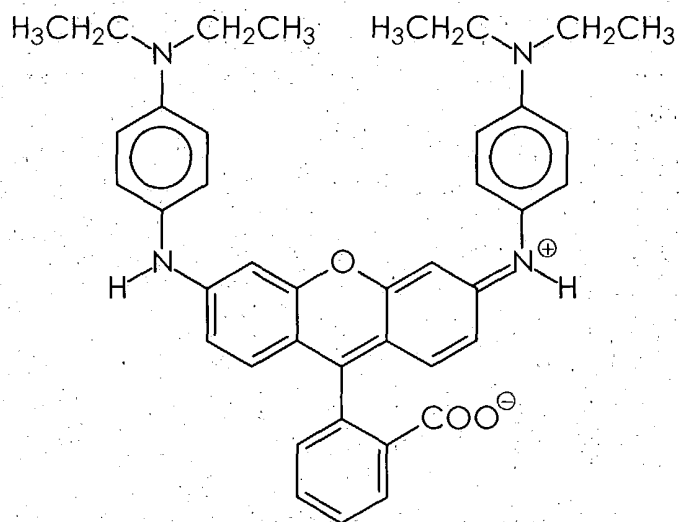


The ring-opened, or protonated, or free-base form of this colorant is
10 believed to be of the formula



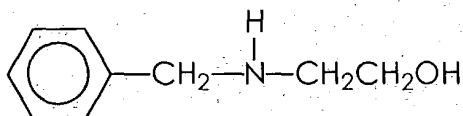
wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula



EXAMPLE XIB

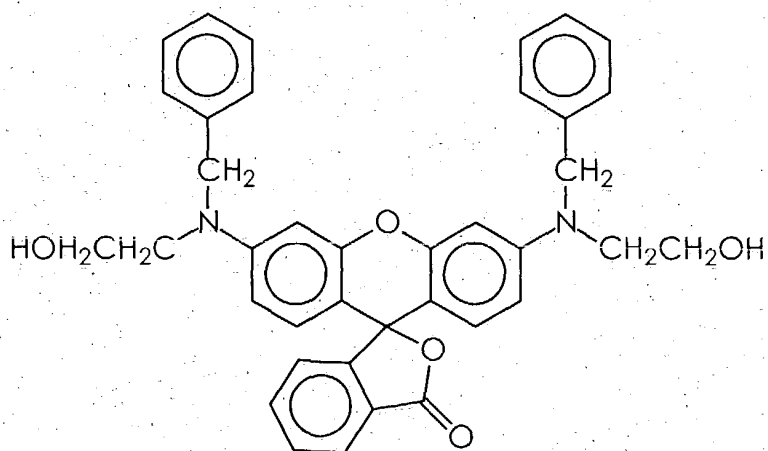
The process of Example IB was repeated except that N-benzylethanolamine (obtained from Aldrich Chemical Co., Milwaukee, WI), of the formula



5 was used instead of distearyl amine. The N-benzylethanolamine was present in an amount of 2.5 moles of N-benzylethanolamine per every one mole of dichlorofluorescein. In addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1 mole of calcium oxide was used per every one mole of dichlorofluorescein, the
10 solvent was dimethyl formamide instead of tetramethylene sulfone, and the reaction mixture was heated to 150°C for 48 hours.

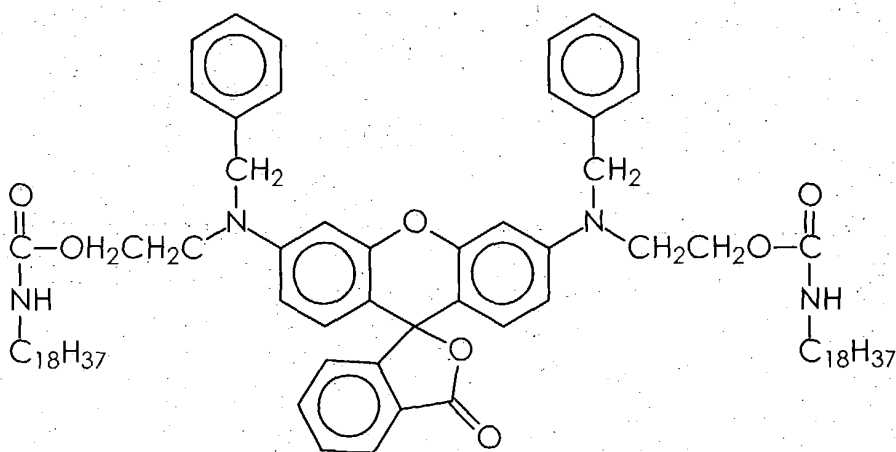
EXAMPLE XIC

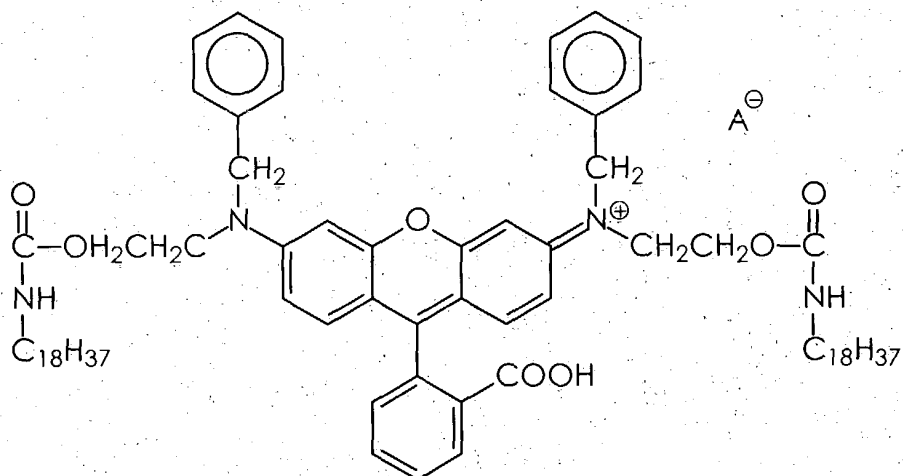
15 The process of Example IC was repeated using the product obtained in Example XIB except that the product was poured into methanol and sufficient EDTA was added to remove all of the Zn²⁺ and Ca²⁺ ions. It is believed that the purified product was of the formula



EXAMPLE XIC-1

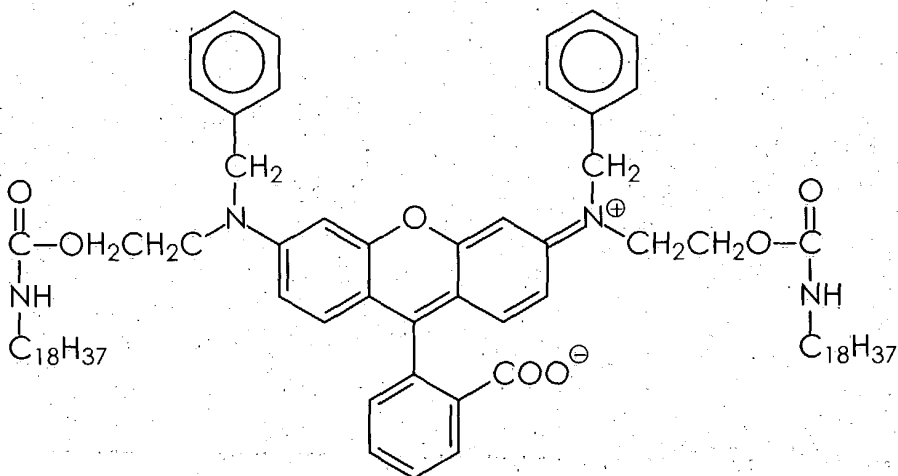
About 10 grams of the product obtained in Example XIC is
 5 added to 9.9 grams of octadecylisocyanate (available from Aldrich
 Chemical Co., Milwaukee, WI) at 120°C, after which 1 drop of
 dibutyltin dilaurate catalyst (available from Aldrich Chemical Co.) is
 added and the reaction is stirred and heated until disappearance of
 the isocyanate peak in the IR is observed. The diurethane rhodamine is
 10 poured into aluminum tins and is believed to be of the formula





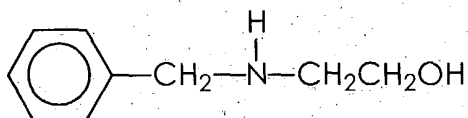
wherein A is the anion corresponding to the acid used for protonation.

5. The zwitterionic form of this colorant is believed to be of the formula



EXAMPLE XIIB

The process of Example IB was repeated except that N-benzylethanolamine (obtained from Aldrich Chemical Co., Milwaukee, WI), of the formula



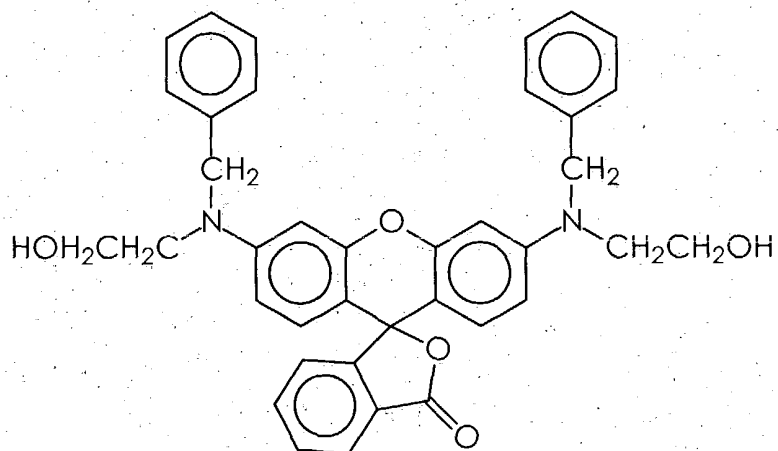
5

was used instead of distearyl amine. The N-benzylethanolamine was present in an amount of 10 moles of N-benzylethanolamine per every one mole of dichlorofluorescein. In addition, 2 moles of zinc chloride were used per every one mole of dichlorofluorescein and 1 mole of calcium oxide was used per every one mole of dichlorofluorescein, the solvent was the excess N-benzylethanolamine instead of tetramethylene sulfone, and the reaction mixture was refluxed in an oil bath for 48 hours, followed by distilling off the excess amine.

15

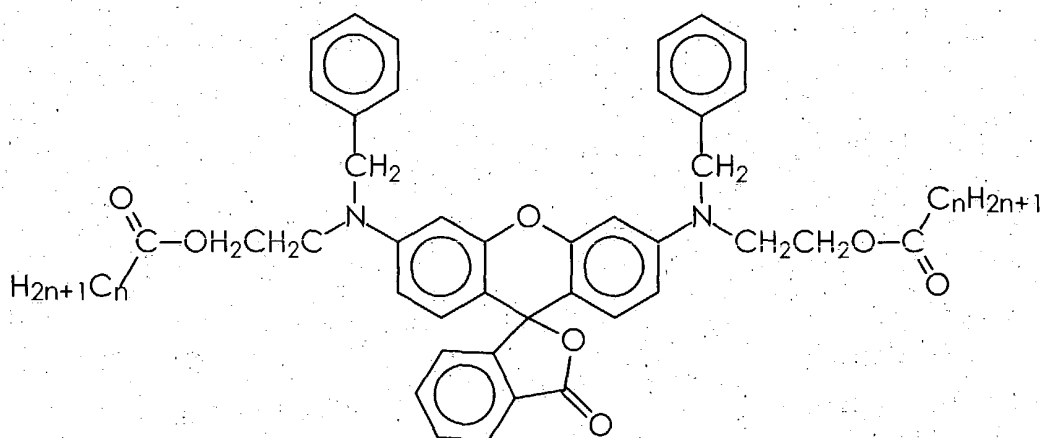
EXAMPLE XIIC

The process of Example IC was repeated using the product obtained in Example XIIB except that the product was poured into methanol and sufficient EDTA was added to remove all of the Zn^{2+} and Ca^{2+} ions. It is believed that the purified product was of the formula

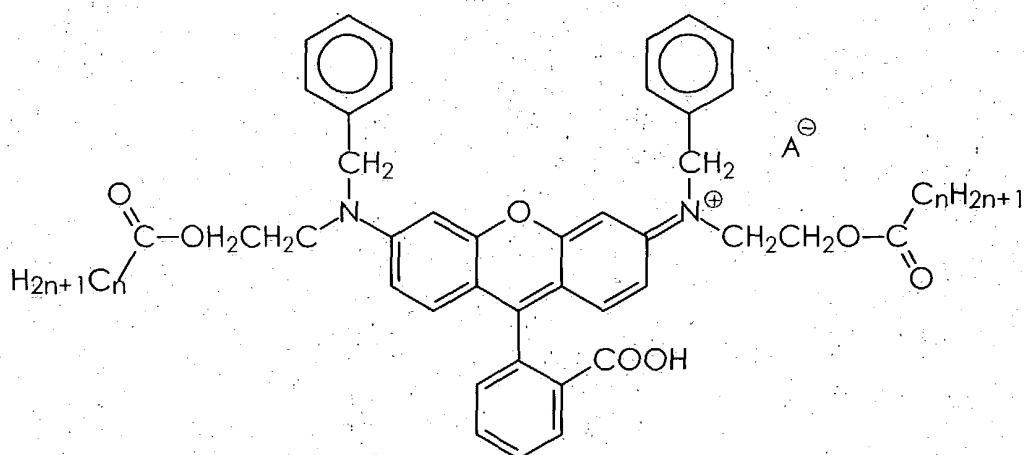


EXAMPLE XIIC-1

In a glass reaction flask is combined 10 grams of the product obtained in Example XIIC, 29.8 grams of UNICID® 700 (a material containing carboxylic acid of the formula RCOOH wherein R is a linear alkyl group having an average of about 50 carbon atoms, also containing other unfunctionalized wax materials in an amount of up to about 25 percent by weight; available from Baker Petrolite, Sugarland, TX), 152 grams of xylene (available from Tarr, Inc., Portland, OR), and 0.6 grams of para-toluenesulfonic acid (available from Capital Resin Corp., Columbus, OH). The materials are mixed and heated to a reflux temperature of about 143°C. After about 72 hours, the reaction is complete. The reaction mixture is then cooled to 40°C and filtered. The filter cake is reslurried and filtered two more times in methanol to remove residual xylene. The filter cake is then dried in air at ambient temperature. It is believed that this filter cake will contain a colorant of the formula



wherein n has an average value of about 50. The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula

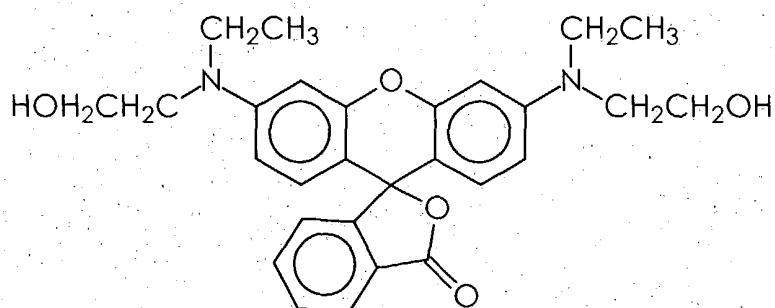


5

wherein A is the anion corresponding to the acid used for protonation. The zwitterionic form of this colorant is believed to be of the formula

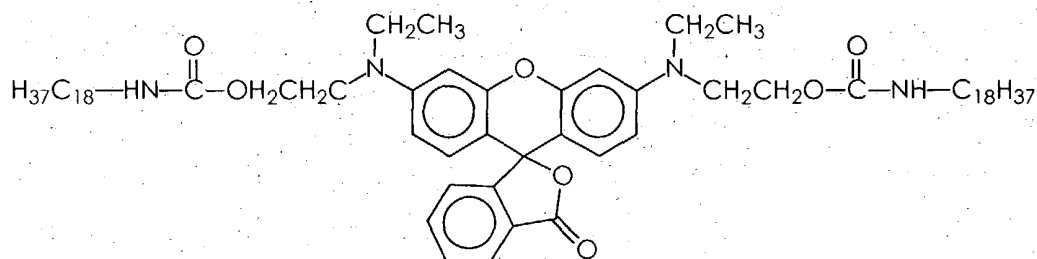
EXAMPLE XIIIIC

The process of Example IC was repeated using the product obtained in Example XIIIIB except that the product was poured into methanol and sufficient EDTA was added to remove all of the Zn^{2+} and Ca^{2+} ions. It is believed that the purified product was of the formula

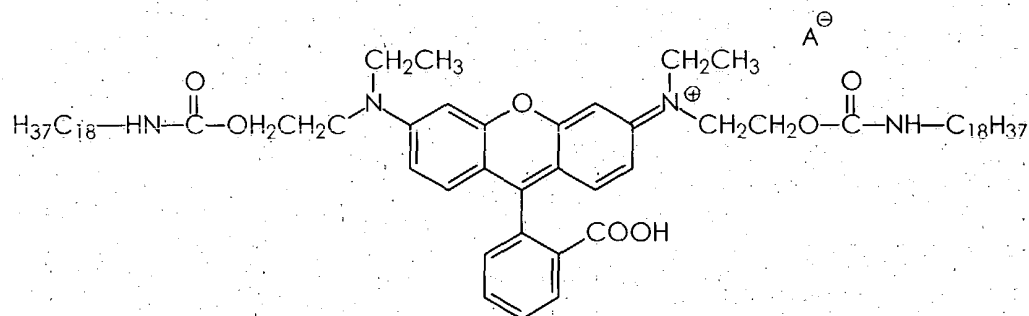


EXAMPLE XIIIIC-1

About 10 grams of the product obtained in Example XIIIIC is added to 12.5 grams of octadecylisocyanate (available from Aldrich Chemical Co., Milwaukee, WI) at 120°C , after which 1 drop of dibutyltindilaurate catalyst (available from Aldrich Chemical Co.) is added and the reaction is stirred and heated until disappearance of the isocyanate peak in the IR is observed. The diurethane rhodamine is poured into aluminum tins and is believed to be of the formula

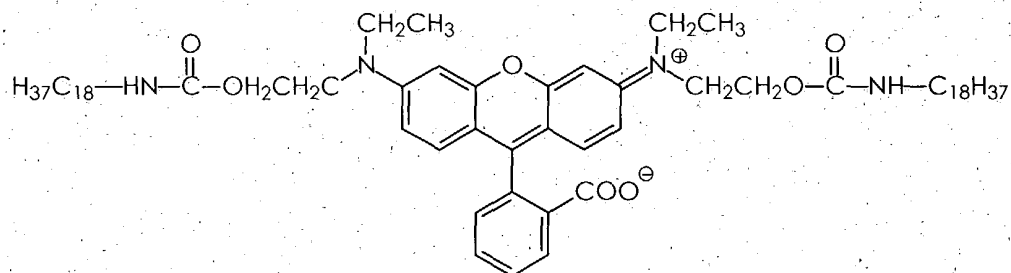


The ring-opened, or protonated, or free-base form of this colorant is believed to be of the formula



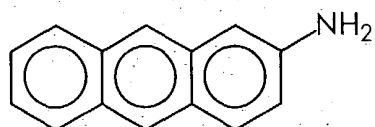
wherein A is the anion corresponding to the acid used for protonation.

5 The zwitterionic form of this colorant is believed to be of the formula



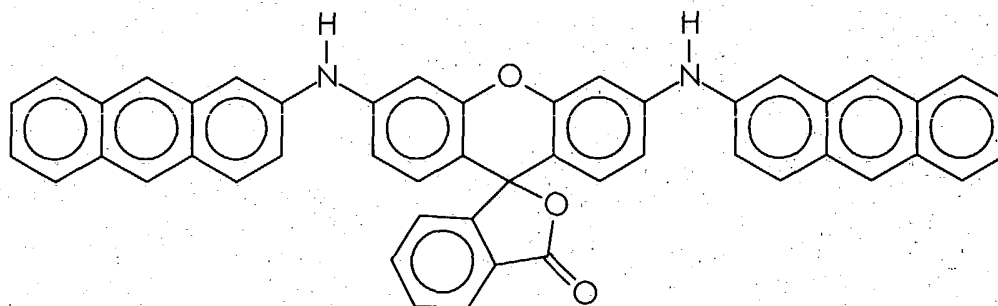
EXAMPLE XIVB

The process of Example IB was repeated except that 2-aminoanthracene (obtained from Aldrich Chemical Co., Milwaukee, WI), of the formula

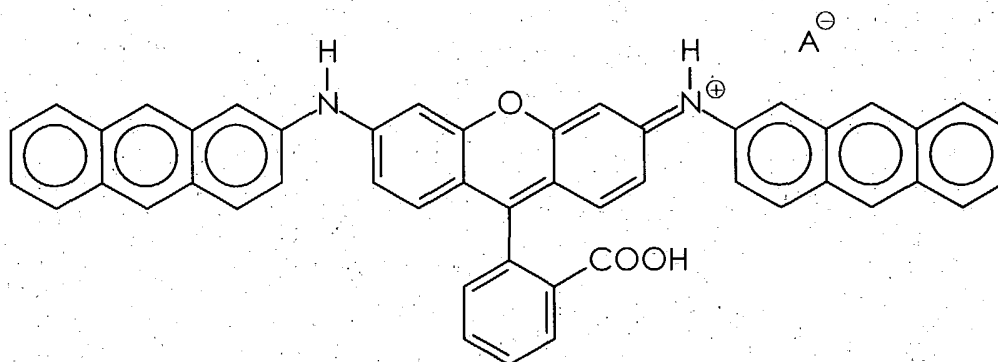


was used instead of distearyl amine. The 2-aminoanthracene was present in an amount of 2.05 moles of 2-aminoanthracene per every

one mole of dichlorofluorescein. It is believed that the product was of the formula

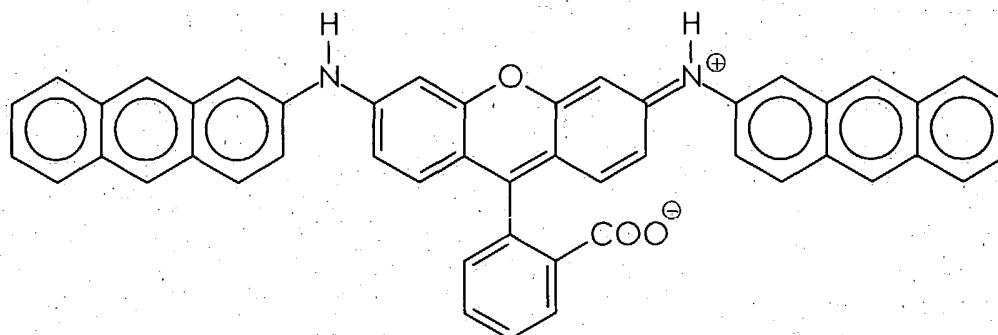


The ring-opened, or protonated, or free-base form of this colorant is
5 believed to be of the formula



wherein A is the anion corresponding to the acid used for protonation.

The zwitterionic form of this colorant is believed to be of the formula

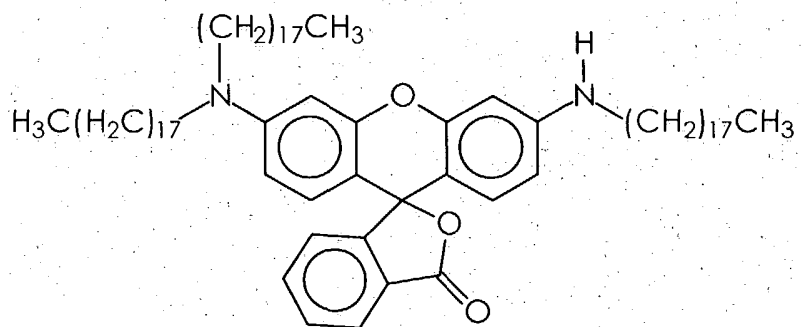
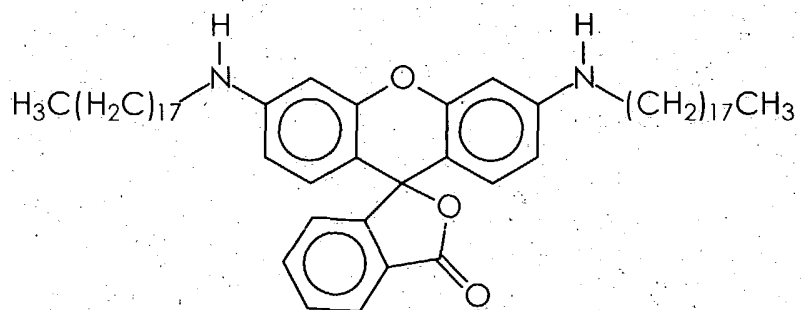


EXAMPLE XVB

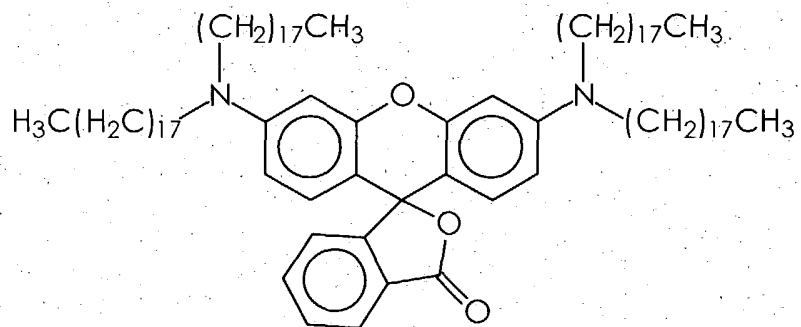
The process of Example IB was repeated except that a
 5 mixture of stearyl amine (ARMEEN 18D; obtained from Akzo-Nobel, McCook, IL) and distearyl amine was used instead of pure distearyl amine. The stearyl amine was present in an amount of 1.02 moles of stearyl amine per every one mole of dichlorofluorescein, and the distearyl amine was present in an amount of 1.02 moles of distearyl
 10 amine per every one mole of dichlorofluorescein.

EXAMPLE XVC

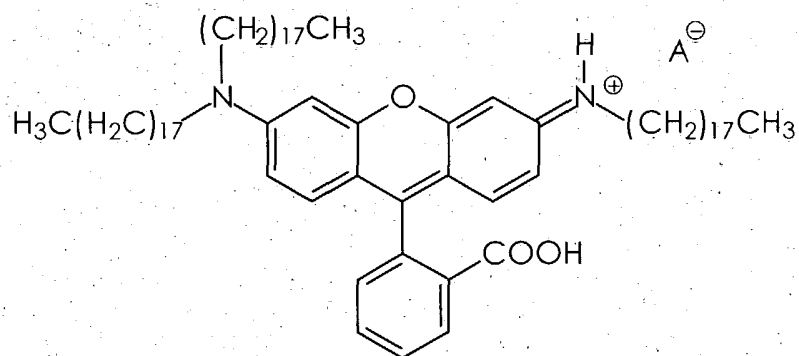
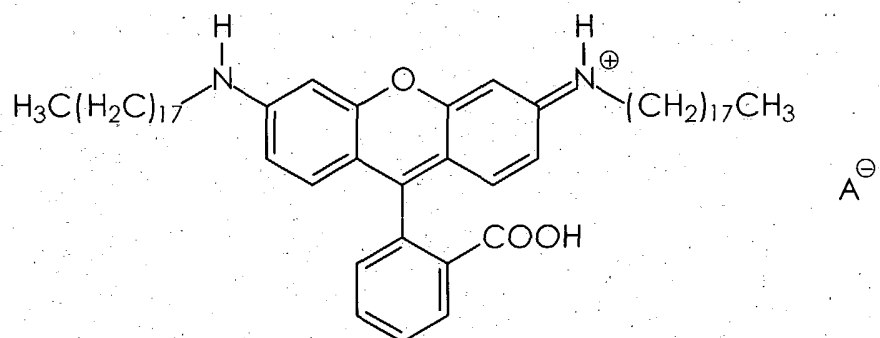
The process of Example IC was repeated using the product obtained in Example XVB. It is believed that the purified product was a
 15 mixture of compounds of the formulae



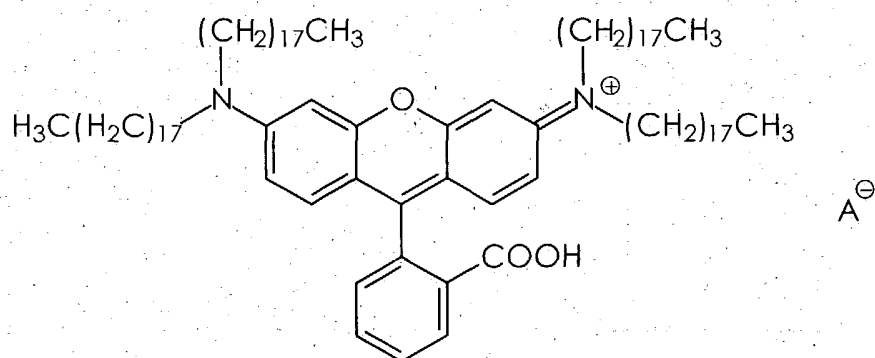
and



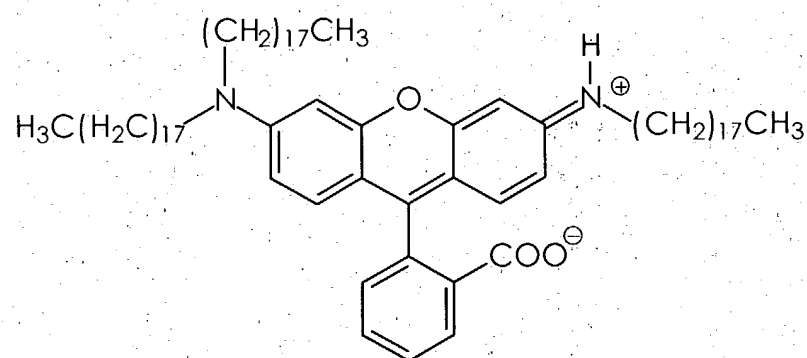
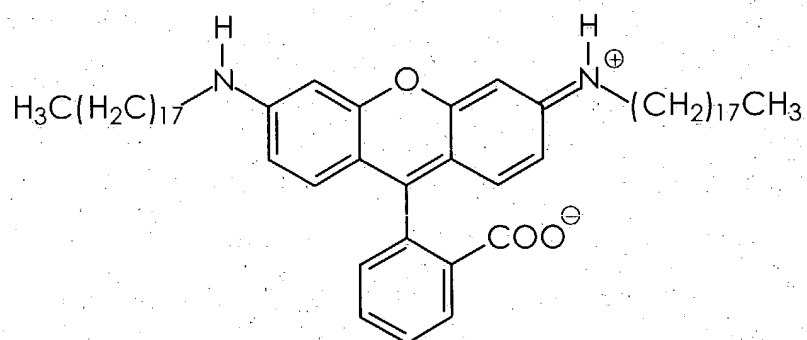
- 5 The ring-opened, or protonated, or free-base forms of these colorants are believed to be of the formulae, respectively,



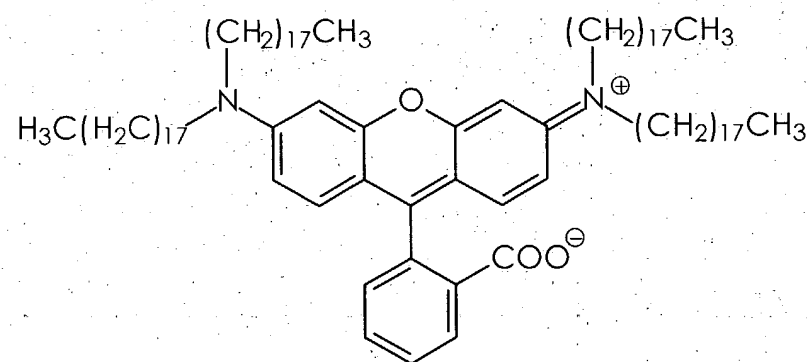
and



- 5 wherein A is the anion corresponding to the acid used for protonation. The zwitterionic forms of these colorants are believed to be of the formulae, respectively,



and



EXAMPLE XVD

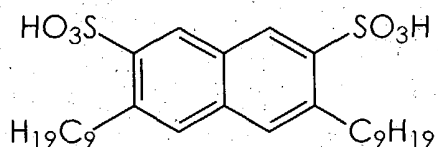
The process of Example ID was repeated using the product obtained in Example XVC.

5

EXAMPLE XVI

The processes of Examples IA through IC were repeated. Thereafter, to the solution of the ring-closed purified tetrastearyl colorant in MIBK was added a naphthalene disulfonate adduct of the formula

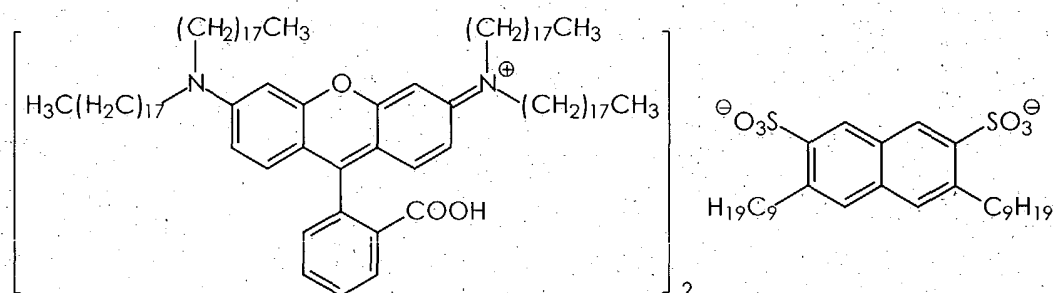
10



15

(dinonylnaphthalene disulfonic acid, 50 wt.% in isobutanol, NACURE® 155, obtained from King Industries, Norwalk, CT) in a stoichiometric amount of 2 moles of naphthalene sulfonate adduct per every one mole of tetrastearyl colorant. The solution was stirred until a magenta color developed fully. Thereafter, the solution was transferred to a 2 liter round bottom flask equipped with distillation setup, and the MIBK was distilled off. The product, a slightly viscous wax when warm, was transferred to a jar and allowed to harden. At room temperature, the product was a deep magenta/red colored somewhat hard wax, believed to be of the formula

20



Ink Preparation and Testing

EXAMPLE XVII

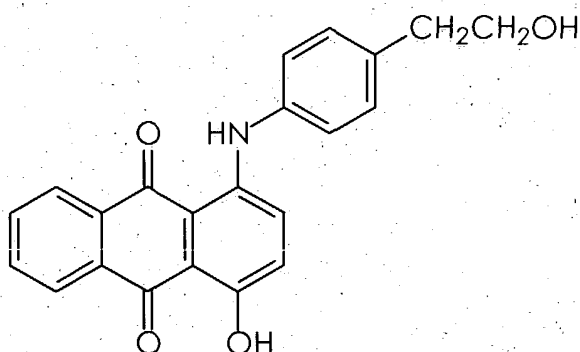
Preparation of Secondary Colorant

PART I

A secondary magenta colorant was prepared as follows.

In a glass reaction flask were combined 73 grams of sublimed quinizarin (obtained from Aceto Corp., Lake Success, NY), 49 grams of leucoquinizarin (obtained from Aceto Corp.), 66 grams of 4-aminobenzene ethanol (obtained from Aceto Corp.), 31 grams of boric acid (obtained from Aldrich Chemical Co., Milwaukee, WI), and 780 grams of methanol (obtained from JT Baker, Phillipsburg, NJ). The materials were mixed and heated until the solvent refluxed at about 66°C.

After about 16 hours of reflux the reaction was complete, having generated an alcohol-substituted colorant of the formula

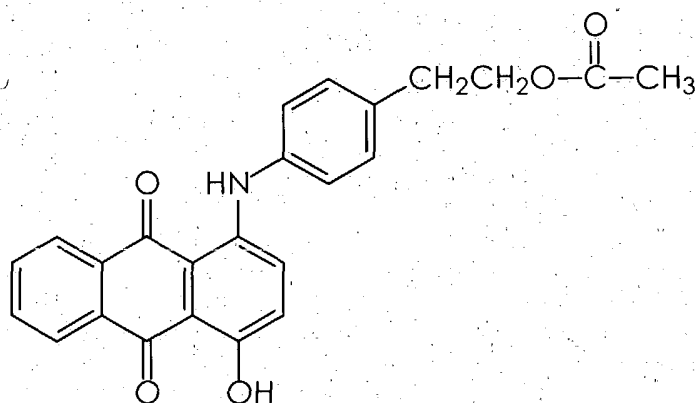


The reaction mixture was cooled and filtered. The product filter cake was dried in air at ambient temperature.

The spectral strength of the alcohol-substituted colorant was determined using a spectrophotographic procedure based on the measurement of the colorant in solution by dissolving the colorant in toluene and measuring the absorbance using a Perkin Elmer Lambda 2S UV/VIS spectrophotometer. The spectral strength of the alcohol-substituted colorant was measured as about 21,000 mL Absorbance Units per gram at absorption λ_{\max} , indicating a purity of about 80 percent.

PART 2

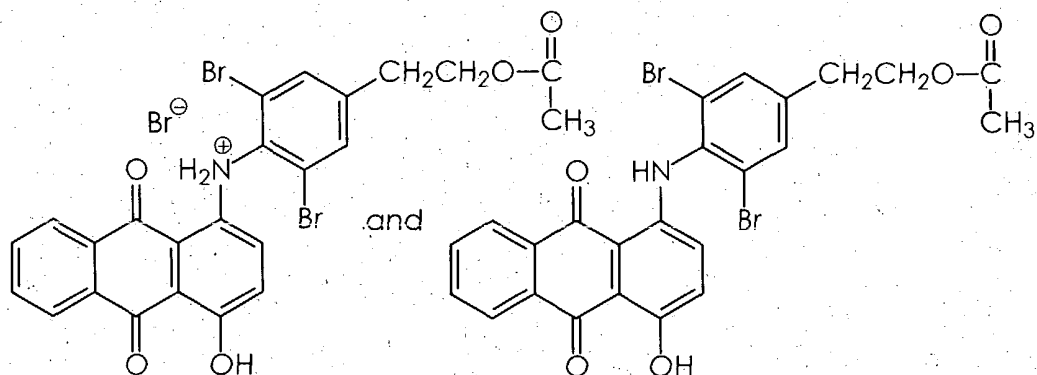
In a glass reaction flask were combined 8 grams of the alcohol-substituted colorant prepared in Part 1 of this Example, 68 grams of glacial acetic acid (obtained from JT Baker), 13 grams of propionic acid (obtained from Aldrich Chemical Co.), and 2.3 grams of acetic anhydride (obtained from Aldrich Chemical Co.). The materials were mixed and heated to a reflux temperature of about 121°C. After about 4 hours of reflux, the reaction was complete and the reaction mixture contained an ethyl acetate-substituted colorant of the formula



PART 3

About 91 grams of the reaction mixture containing the ethyl acetate-substituted colorant from Part 2 of this Example was charged into a glass reaction flask. The mixture was cooled to a minimum of 30°C. While mixing, about 9 grams of bromine (obtained from Aldrich Chemical Co.) was added to the mixture at a rate such that the temperature remained below about 40°C. The mixture was then heated to about 40°C. After about 24 hours of mixing the reaction was complete.

The reaction mixture was then quenched into 234 grams of deionized water and allowed to cool to room temperature. The reaction mixture was then filtered. The filter cake was reslurried and filtered twice in deionized water to remove most of the residual acetic acid. The filter cake was then dried in a 60°C oven. This filter cake contained a mixture of brominated ethyl acetate-substituted colorants of the formulae



The spectral strength of the brominated ethyl acetate-substituted colorant was determined using a spectrophotographic procedure based on the measurement of the colorant in solution by dissolving the colorant in toluene and measuring the absorbance using a Perkin Elmer Lambda 2S UV/VIS spectrophotometer. The spectral strength of the brominated ethyl acetate-substituted colorant was measured as about 15,000 mL Absorbance Units per gram at absorption λ_{max} . This spectral strength indicated a purity of about 60 percent.

10

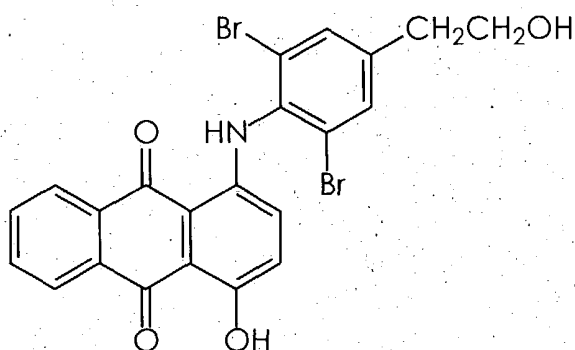
PART 4

In a glass reaction flask were combined 18 grams of the mixture of the brominated ethyl acetate-substituted colorant and its salt prepared in Part 3 of this Example, 72 grams of N-methyl-2-pyrrolidone (obtained from Aldrich Chemical Co.), 4 grams of sodium hydroxide (obtained from Aldrich Chemical Co.), and 4 grams of deionized water. The materials were mixed and heated to about 60°C. After about 3 hours the reaction was complete.

The reaction mixture was then quenched into 234 grams of deionized water and allowed to cool to room temperature. Glacial

20

acetic acid was added until the solution reached a pH of between 6 and 7. The reaction mixture was then filtered. The filter cake was reslurried and filtered twice in deionized water to remove most of the residual N-methyl-2-pyrrolidone. The filter cake was then dried in a 60°C oven. This filter cake contained a brominated alcohol-substituted colorant of the formula



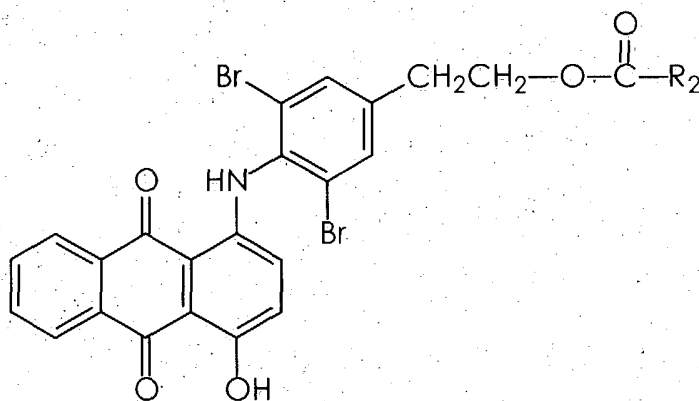
The spectral strength of the brominated alcohol-substituted colorant was determined using a spectrophotographic procedure based on the measurement of the colorant in solution by dissolving the colorant in an equal mixture of toluene and tetrahydrofuran and measuring the absorbance using a Perkin Elmer Lambda 2S UV/VIS spectrophotometer. The spectral strength of the brominated alcohol-substituted colorant was measured as about 16,000 mL Absorbance Units per gram at absorption λ_{\max} . This spectral strength indicated a purity of about 60 percent.

PART 5

In a glass reaction flask were combined 16 grams of the brominated alcohol-substituted colorant prepared in Part 4 of this Example, 31 grams of UNICID® 700 (a material containing carboxylic

acid of the formula R_2COOH wherein R_2 is a linear alkyl group having an average of about 50 carbon atoms, also containing other unfunctionalized wax materials in an amount of up to about 25 percent by weight; obtained from Baker Petrolite, Sugarland, TX), 152 grams of xylene (obtained from Tarr, Inc., Portland, OR), and 0.6 grams of para-
5 toluenesulfonic acid (obtained from Capital Resin Corp., Columbus, OH). The materials were mixed and heated to a reflux temperature of about 143°C . After about 7 hours, the reaction was complete.

The reaction mixture was then cooled to 40°C and filtered.
10 The filter cake was reslurried and filtered two more times in methanol to remove residual xylene. The filter cake was then dried in air at ambient temperature. This filter cake contained a colorant of the formula



wherein R_2 is a linear alkyl group having an average of about 50
15 carbon atoms.

The spectral strength of the colorant was determined using a spectrophotographic procedure based on the measurement of the colorant in solution by dissolving the colorant in an equal mixture of toluene and tetrahydrofuran and measuring the absorbance using a
20 Perkin Elmer Lambda 2S UV/VIS spectrophotometer. The spectral strength of the colorant was measured as about 5,000 mL Absorbance

Units per gram at absorption λ_{\max} . This spectral strength indicated a purity of about 40 percent.

Ink compositions containing the colorants of Examples ID
5 and IIID, and, for comparison purposes, commercially available n-butyl Solvent Red 172 (n-BuSR172; UNIGRAPH Red 1900, obtained from United Color Manufacturing, Inc., Newtown, PA) and commercially available Solvent Red 49 (SR49; a rhodamine colorant obtained from BASF, Germany) were prepared as follows.

10 Ink A: In a stainless steel beaker were combined 243.00 grams of polyethylene wax (PE 655, obtained from Baker Petrolite, Tulsa, OK, of the formula $\text{CH}_3(\text{CH}_2)_{50}\text{CH}_3$), 122.15 grams of stearyl stearamide wax (KEMAMIDE® S-180, obtained from Crompton Corporation, Greenwich, CT), 108.00 grams of a tetra-amide resin obtained from the
15 reaction of one equivalent of dimer diacid with two equivalents of ethylene diamine and UNICID® 700 (a carboxylic acid derivative of a long chain alcohol obtained from Baker Petrolite, Tulsa, OK), prepared as described in Example 1 of U.S. Patent 6,174,937, the disclosure of which is totally incorporated herein by reference, 65.45 grams of a
20 urethane resin obtained from the reaction of two equivalents of ABITOL® E hydroabietyl alcohol (obtained from Hercules Inc., Wilmington, DE) and one equivalent of isophorone diisocyanate, prepared as described in Example 1 of U.S. Patent 5,782,966, the disclosure of which is totally incorporated herein by reference, 23.00
25 grams of a urethane resin that was the adduct of three equivalents of stearyl isocyanate and a glycerol-based alcohol, prepared as described in Example 4 of U.S. Patent 6,309,453, the disclosure of which

is totally incorporated herein by reference, and 1.10 gram of NAUGUARD® 445 antioxidant (obtained from Uniroyal Chemical Co., Middlebury, CT). The materials were melted together at a temperature of 135°C in an oven, and then blended by stirring in a temperature-controlled mantle at 135°C for 0.2 hour. To this mixture was then added 22.98 grams of the tetrastearyl colorant prepared as described in Example ID, 17.66 grams of a secondary magenta colorant (prepared as described in Parts 1 through 5 of this Example), and 3.20 grams dodecyl benzene sulfuric acid (DDBSA, Bio-soft S-100, obtained from Stepan Company, Elwood, IL). After stirring for 2 additional hours, the magenta ink thus formed was filtered through a heated MOTT® apparatus (obtained from Mott Metallurgical) using Whatman #3 filter paper under a pressure of 15 pounds per square inch. The filtered phase change ink was poured into molds and allowed to solidify to form ink sticks. The magenta phase change ink thus prepared exhibited a viscosity of 10.49 centipoise as measured by a Rheometrics cone-plate viscometer at about 140°C, melting points of 85°C and 103°C as measured by differential scanning calorimetry using a DuPont 2100 calorimeter, a glass transition temperature (T_g) of 19°C, and a spectral strength of 1,023 milliliters absorbance per gram at 552 nanometers, determined by using a spectrophotographic procedure based on the measurement of the colorant in solution by dissolving the solid ink in n-butanol and measuring the absorbance using a Perkin Elmer Lambda 2S UV/VIS spectrophotometer.

Ink B: Ink B was prepared in a similar manner to that used to prepare Ink A but using a different formulation for the ink composition as described in the table below. The properties of Ink B were obtained

using the same methods as those used for Ink A. As shown in the table, the predominant difference between Ink A and Ink B is the relative concentrations of colorants and DDBSA in the ink. The viscosities of Ink A and Ink B are virtually the same.

5 Ink C: Ink C was prepared by the process described for Ink A except that the distearyl colorant from Example IIID was used in place of the tetrastearyl colorant from Example ID and no DDBSA was used. The properties of Ink C were obtained using the same methods as those used for Ink A.

10 Inks D and E: These inks were prepared by the process described for Ink A except that the distearyl colorant from Example IIID was used in place of the tetrastearyl colorant from Example ID and no DDBSA was used. The properties of Inks D and E were obtained using the same methods as those used for Ink A. Inks D and E use same
15 colorant set as in Ink C but have higher relative concentrations of the distearyl colorant, as the results in high spectra strength indicate.

 Ink F (Comparative): An ink was prepared by the process described for Ink A except that instead of the colorant from Example ID and the secondary magenta colorant prepared in Parts 1 through 5 of
20 this Example, the commercially available SR 49 and n-butyl SR172 were used. The properties of Ink F were obtained using the same methods as those used for Ink A.

 Ink G (Comparative): An ink was prepared by the process described for Ink A except that instead of the colorant from Example
25 ID, the commercially available SR 49 was used. The properties of Ink G were obtained using the same methods as those used for Ink A.

Ink H: An ink was prepared by the process described for Ink A except that instead of the colorant from Example ID, the colorant prepared in Example XVI was used, dinonylnaphthalene disulfonic acid (50 wt.% in isobutanol, NACURE® 155, obtained from King Industries, Norwalk, CT) was used in place of DDBSA, and no secondary colorant was used. The properties of Ink H were obtained using the same methods as those used for Ink A.

The following table summarizes the compositions of the various inks and the amounts of ingredients (weight percentage numbers given in the table) therein:

Ingredient	Ink A	Ink B	Ink C	Ink D	Ink E	Ink F	Ink G	Ink H
POLYWAX	40.06	40.16	44.62	43.81	42.99	41.83	45.67	38.68
S-180	20.14	19.38	13.18	13.18	12.93	19.26	13.17	17.05
Tetra-amide	17.81	17.82	19.36	19.00	18.64	20.36	19.04	23.56
Urethane Resin 1*	10.79	12.47	10.42	10.42	10.22	10.13	10.68	10.45
Urethane Resin 2**	3.79	4.26	8.00	7.90	7.75	5.56	8.09	5.76
Example ID colorant	3.79	2.61	---	---	---	---	---	---
Example IIID colorant	---	---	2.30	3.57	5.33	---	---	---
Example XVI colorant	---	---	---	---	---	---	---	3.56
SR49	---	---	---	---	---	0.55	0.46	---
2°magenta colorant	2.91	2.03	1.93	1.93	1.93	---	1.91	---
n-Butyl SR 172	---	---	---	---	---	1.01	---	---
DDBSA	0.53	1.10	---	---	---	1.11	0.80	---
NACURE 155	---	---	---	---	---	---	---	0.78
NAUGUARD 445	0.18	0.18	0.20	0.20	0.20	0.19	0.20	0.18
Total	100.0	100.0	100.0	100.0	100.0	100.0	100.0	100.0

* ABITOL E based urethane resin

** Glycerol alcohol based urethane resin

The magenta inks thus prepared were successfully printed on HAMMERMILL LASERPRINT® paper (obtained from International Paper, Memphis, TN) with a XEROX® PHASER 860 printer, which uses a printing process wherein the ink is first jetted in an imagewise pattern onto an intermediate transfer member followed by transfer of the imagewise pattern from the intermediate transfer member to a final recording substrate. The solid field images with a resolution of 450 dpi x 600 dpi were generated from the printer, and their color space data were obtained on an ACS® Spectro Sensor® II Colorimeter (obtained from Applied Color Systems Inc.) in accordance with the measuring methods stipulated in ASTM 1E805 (Standard Practice of Instrumental Methods of Color or Color Difference Measurements of Materials) using the appropriate calibration standards supplied by the instrument manufacturer. For purposes of verifying and quantifying the overall colorimetric performance of the inks, measurement data were reduced, via tristimulus integration, following ASTM E308 (Standard Method for Computing the Colors of Objects using the CIE System) in order to calculate the 1976 CIE L* (Lightness), a* (redness-greenness), and b* (yellowness-blueness) CIELAB values for each phase change ink sample.

Another type of printed sample was generated on HAMMERMILL LASERPRINT® paper using a K Printing Proofer (manufactured by RK Print Coat Instrument Ltd., Litlington, Royston, Heris, SG8 0OZ, U.K.). In this method, the tested inks were melted onto a printing plate set at 150°C temperature. A roller bar fitted with the paper was then rolled over the plate containing the melted ink on its

surface. The ink on the paper was cooled, resulting in three separated images of rectangular blocks. The most intensely colored block contained the most ink deposited on the paper, and was therefore used to obtain the color value measurements.

- 5 Printed samples of the magenta inks both from the XEROX PHASER® printer and from the K-Proofer were evaluated for color characteristics, which are reported in the table below. As is apparent, the CIE L*a*b* values for inks made with colorants according to the present invention represent a magenta shade printed ink. The table
- 10 below lists the viscosity (η , centipoise) of the inks at 140°C, the spectral strength in n-butanol (SS, mL*g⁻¹cm⁻¹) and absorbance maximum (Lambda max, nm) of the inks; and the CIE L*a*b color coordinates of the prints made either using the XEROX PHASER® 860 printer or the K-proofer:

15

Ink	η	SS	Lambda Max	L*/a*/b* of Prints from 860 Printer	L*/a*/b* of prints from K-Proofer
A	10.49	10.23	552	48.7/76.9/-37.7	---
B	10.54	1328	552	50.1/69.1/-37.2	56.3/59.3/-32.5
C	10.62	1382	522	---	62.2/71.2/-29.9
D	10.30	2075	522	---	58.6/75.0/-27.8
E	10.59	2874	522	---	54.8/78.0/-25.2
F	10.53	1686	555	---	---
G	10.77	1279	555	54.0/76.8/-41.3	60.9/68.0/-42.7
H	---	---	---	---	65.3/70.5/-30.0

--- indicates not measured

The color values in the above table indicate that the colorant of Example ID, the colorant of Example IIID, and the colorant of Example XVI can be used in hot melt inks with good magenta color as evidenced by the high a^* and b^* values of the prints. As evidenced in the table, the colors of the inks containing the tetrastearyl colorant (Ink A and B) of this invention are bright magenta and similar to the comparative inks made from commercially available SR 49 (Ink G). In contrast to commercial SR 49 colorant, which normally needs a relatively strong acid such as DDBSA to develop its color in ink base, the distearyl colorants of this invention in inks C through E show reasonably strong reddish magenta color without an acid developer. The inks' spectral strengths and a^* values increase with their colorant concentration, as shown in Inks C through E. Good colorant solubility of the distearyl colorant in tested ink bases are demonstrated by the very high colorant loads and corresponding very high spectral strength in Ink C through E.

EXAMPLE XVIII

Thermal stability of the tetrastearyl colorant from Example ID was compared to comparative SR 49 colorant by monitoring their inks' spectral strength after heating. In this method, the inks were heated in glass jars continuously in an oven of 145°C. The ink's spectral strength in n-butanol (related to the amount of the soluble colorant in ink) was measured as a function of aging time. Any thermal decomposition of a test colorant would be observable as discoloration and could thereby detected as a change in the spectral strength.

The table below illustrates spectral strength relative to the initial values verse elapsed time for Ink A and comparative Ink F samples. The remaining spectral strength is used here as the measurement of remaining undegraded colorant in the ink. It is evident
5 that the colorants in Ink A have superior thermal stability compared to those in comparative Ink F.

Cooking time (days)	1	2	7	11	15
Ink A	100.0	97.1	86.9	79.4	72.7
Ink F	100.0	94.1	80.0	71.1	63.7

Colorant degradation not only leads to a decrease in
10 spectral strength (as shown in the above table) but can also generate undesirable color shift or fade as a result of the colorant decomposition reaction in the ink. Both of these phenomena can adversely affect the color quality of prints from the inks if the colorant is not thermally stable. From a practical application point of view, the overall color change of
15 the ink (measured as ΔE change in color values) is also important when evaluating colorant thermal stability.

A thermal stability test was performed by heating the test inks in glass jars, then sampling and printing the inks on HAMMERMILL LASERPRINT® paper using a K-Proofer, and finally measuring the color
20 changes of the prints of the sampled inks as a function of time. The color changes of the resultant prints were monitored by CIELAB values and expressed by Delta E relative to the initial CIELAB values. The color change of each sample was determined according to the methods described hereinabove for obtaining CIELAB values. Color changes

were determined following ASTM D2244-89 (Standard Test Method for Calculation of Color Differences From instrumentally Measured Color Coordinates) ($\Delta E = [(L^*_1 - L^*_2)^2 + (a^*_1 - a^*_2)^2 + (b^*_1 - b^*_2)^2]^{1/2}$). The results for Inks B, C, and G are shown in the tables below. As the data in the tables indicate, Inks B and C containing the colorants of Example ID and Example IIID demonstrated better color stability than comparative Ink G containing SR 49.

Cooking time (days)	0	1	3	5	7	9
Ink C	0.0	2.5	4.3	3.6	8.3	10.2
Ink G	0.0	4.2	8.5	10.0	11.5	11.7

ΔE values for Inks C and G when heated at 140°C

Cooking time (days)	0	0.8	1.0	3.0	3.1	5.0	6.9	7.0	10.0
Ink B	0.0	---	1.8	3.6	---	5.0	---	5.3	---
Ink G	0.0	2.6	---	---	4.8	---	8.7	---	9.4

ΔE values for Inks B and G when heated at 136°C; --- indicates not measured.

EXAMPLE XIX

The ink containing the colorant of Example ID (Ink B) was tested for colorant diffusion. For comparison purposes, comparative Ink G containing the commercial colorant SR 49 was also tested; both inks contained the secondary colorant prepared in Parts 1 through 5 of

Example XVII and DDBSA. This diffusion evaluation method entailed generating text prints of the magenta inks, applying SCOTCH® tape (obtained from 3M, St. Paul, MN) adhesive to a portion of the text, and examining the extent of colorant diffusion into the adhesive material over time. The text print samples were then studied to observe how heat amplified colorant diffusion over time. This test simulated the real situation when one applies SCOTCH® tape adhesive labels onto prints generated with inks containing the colorants. Heating the prints served to mimic the conditions when prints are handled in warm climates or passed through high-speed document handlers.

After heating at 45°C for six days and then remaining at room temperature for six weeks, the printed text area using comparative Ink G with commercial colorant SR 49 displayed extensive colorant diffusion into the applied tape, resulting in blurry edges of the text characters. For Ink B containing the colorant of Example ID, the extent of diffusion was noticeable but of lesser extent, and the printed text situated beneath the adhesive tape was more legible. The results demonstrated that the colorant of Example ID has fewer tendencies to diffuse compared to the commercial SR 49. Although not being limited to any particular theory, it is believed that the long alkyl groups in the colorant of Example ID help to prevent colorant diffusion in the ink base.

Other embodiments and modifications of the present invention may occur to those of ordinary skill in the art subsequent to a review of the information presented herein; these embodiments and

modifications, as well as equivalents thereof, are also included within the scope of this invention.

The recited order of processing elements or sequences, or the use of numbers, letters, or other designations therefor, is not
5 intended to limit a claimed process to any order except as specified in the claim itself.